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Published in:
Journal of Physics D-Applied Physics

DOI:
[10.1088/0022-3727/43/37/374015](https://doi.org/10.1088/0022-3727/43/37/374015)

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Document Version
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Publication date:
2010

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Citation for published version (APA):

Gengler, R. Y. N., Spyrou, K., & Rudolf, P. (2010). A roadmap to high quality chemically prepared graphene. *Journal of Physics D-Applied Physics*, 43(37), 374015-1-374015-19. [374015].
<https://doi.org/10.1088/0022-3727/43/37/374015>

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A roadmap to high quality chemically prepared graphene

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Received 9 April 2010, in final form 13 May 2010

Published 2 September 2010

Online at stacks.iop.org/JPhysD/43/374015

Abstract

Graphene was discovered half a decade ago and proved the existence of a two-dimensional system which becomes stable as a result of 3D corrugation. It appeared very quickly that this exceptional material had truly outstanding electronic, mechanical, thermal and optical properties. Consequently a broad range of applications appeared, as the graphene science speedily moved forward. Since then, a lot of effort has been devoted not only to the study of graphene but also to its fabrication. Here we review the chemical approaches to graphene production, their advantages as well as their downsides. Our aim is to draw a roadmap of today's most reliable path to high quality graphene via chemical preparation.

(Some figures in this article are in colour only in the electronic version)

In the past few years much interest has focused on graphene, a flat monolayer of carbon atoms tightly packed into a two-dimensional (2D) honeycomb lattice. Novoselov and Geim's discovery, that micromechanical cleavage of bulk graphite allows us to isolate graphene [1], has triggered a tremendous amount of scientific interest for this new material, at first mainly for its electronic properties. Graphene was immediately seen as the successor of the current silicon-based technology since the 2D honeycomb carbon structure features semi-metallic behaviour and high carrier mobilities, which are ideal for a potential implementation as computing element. Therefore, graphene sheets, if processable, could be used in classical computers, even though examples of use as qubit elements or for spintronics potentially extend its use further then to purely classical transistor-based microprocessors. Alongside this main research stream, pure graphene was found to exhibit outstanding mechanical [2], thermal [3] and optical properties [4], while graphene derivatives like graphene oxide or other types of functionalized graphene display remarkable catalytic, mechanical, sensing and electronic properties.

Although graphene has so far been explored mainly in fundamental research, creating or depositing high quality graphene for application purposes has been one of the main challenges. Without realizing the future impact of his research,

Brodie was the first to produce exfoliated graphite (through oxidation) [5]. The final product was not graphene but graphene oxide (noticed by Brodie because of the increased mass of the reaction product) [5]. More than a century later, a drastically different method, micromechanical cleavage [1], used to produce isolated graphene sheets, revealed graphene to the physics world. Since then chemists and physicists have used a plethora of (more or less successful) methods, among which chemical vapour deposition (CVD) is one of the most promising [6] in terms of coverage and layer quality. In fact, while micromechanical cleavage has been very successful in producing samples for fundamental studies, its extremely low yield and lack of control in placement of the sheets give it no future in application. Other methods based on the use of SiC as a substrate and as a precursor have excellent potential in the pursuit of the best quality synthetic graphene [7–11]. An alternative road is chemically prepared graphene, which we review in this contribution, focusing on properties and quality as a function of the production method (electronic properties, transparency, ambipolar behaviour, sheet size and related features such as defect/impurity level and coverage on various substrates).

Chemical preparation processes of graphene sheets use graphite as starting material and delaminate it by various means, namely by chemical functionalization, oxidation or by intercalation. To give a good representation of

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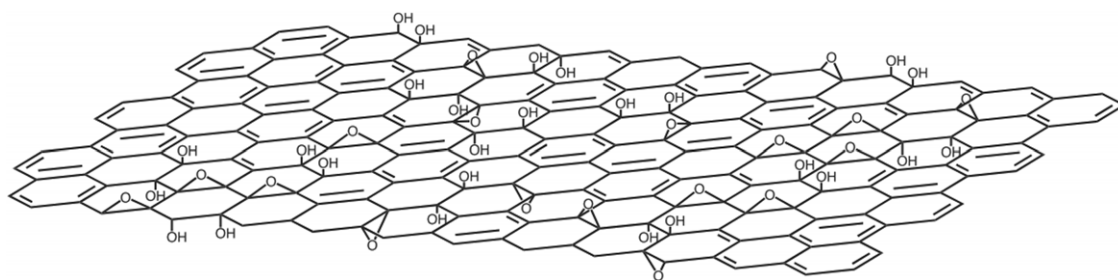


Figure 1. Schematic representation of the structure of graphene oxide sheets (GO); grafted hydroxyl and epoxide groups disrupt the sp^2 -bonded carbon mesh of pure graphene. Scheme taken from Park and Ruoff. Reprinted with permission from Macmillan Publishers Ltd: Nature Nanotechnology [23] Copyright 2009.

today's scientific landscape concerning chemical methods for graphene production we divided the subject into two subcategories, the preponderant graphene oxide (GO) production as a first one and all the other methods (intercalation with N-methylpyrrolidone (NMP), functionalized surfactants, perfluorinated aromatic molecules, etc) as a second one.

1. Graphene from graphene oxide

The first reports on the production of GO date from 1840 by Schaffhaeütl [12] and 1859 with Brodie [5]. In the latter GO was prepared by treating natural graphite (Ceylon) with an oxidation mixture consisting of potassium chlorate and fuming nitric acid, for reaction times of 3–4 days with the solution kept at a temperature of 60 °C, until no further change was observed. Interestingly, the author, Brodie, also proposed an alternative method, where the heating was replaced by exposing the oxidation mixture flasks to sunlight and described it as advantageous because it was faster. Staudenmaier [13] proposed a variation to Brodie's method, where the graphite is oxidized in concentrated sulfuric and nitric acids with potassium chlorate. 99 years after Brodie's first experiments, Hummers and Offeman [14] presented a method where the oxidation of graphite to graphitic oxide is accomplished by treating graphite with a water-free mixture of concentrated sulfuric acid, sodium nitrate and potassium permanganate. This process requires less than 2 h for completion and rather low temperatures (below 45 °C). In the pursuit of high quality graphene these three methods have been used extensively over the past few years. Based on the results obtained with a variety of modern analysis techniques such as NMR [15], XPS [16–18], TEM [19] and Raman [20, 21], the most recent model of GO's structure is the one depicted schematically in figure 1, where hydroxyl and epoxide groups grafted randomly (for older structural models of GO and details of derivation of this model see [22]) to the carbon mesh alter the sp^2 -bonded carbon network of pure graphene sheets [23]. Owing to the presence of such hydrophilic polar groups, the GO presents swelling and intercalation properties very similar to clay [24]. In the next four sections, we shall review some of the characteristics of this new material.

2. Ambipolar behaviour—the quest for high field effect mobility

While an incredibly large number of publications on graphene have appeared in the last 3–4 years, very few actually testify to the truly amazing properties of this material. In particular, most of the recent publications concerning the development of new synthesis methods for graphene, comprising exfoliation, reduction, deposition, etc, omit giving evidence for ambipolar behaviour by recording the Dirac curve, as the conductivity, resistivity or drain–source current versus gate voltage measured in a field effect transistor [FET] device configuration is called. Transmission electron microscopy and atomic force microscopy, often used to demonstrate the graphene character of newly produced material, are very local probes and do not inform on the overall graphene quality. If one wants to compare chemically produced material with high quality graphene created by micromechanical cleavage, the Dirac curve is the best non-local probe. Both graphene and graphite display good conductivity; therefore measuring a high flake conductivity without gate voltage dependence does not prove a bipolar behaviour. To the best of our knowledge, the measurements reported by Gómez-Navarro *et al* [20] were the first to actually show the ambipolar character of the deposited chemically prepared material, i.e. reduced graphene oxide (RGO). The room-temperature field effect mobilities of 2–200 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$ for holes and 0.5–30 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$ for electrons reported for these reduced GO samples are approximately 2 orders of magnitude lower than those of mechanically cleaved graphene [1] and definitely lower than the 200 000 $\text{cm}^2 \text{V}^{-1} \text{s}^{-1}$ field effect mobility of free-standing graphene [25] because of the defective nature of the reduced layers (see section below). The reader should be aware that field effect mobility values depend on channel length and electrode material, therefore the cited numbers (here and below) always refer to the devices employed in the cited works. As we will discuss further, extracting mobility values from incomplete Dirac curves, like the ones recorded by Gómez-Navarro *et al* [20], where the metallic regime is not reached, is a non-trivial task and gives only approximate values. The data, shown in figure 2, collected at various temperatures under a low pressure of helium exhibit maximal resistance close to zero gate voltage because of this inert atmosphere. Prolonged exposure to ambient air (>24 h) resulted in a pronounced

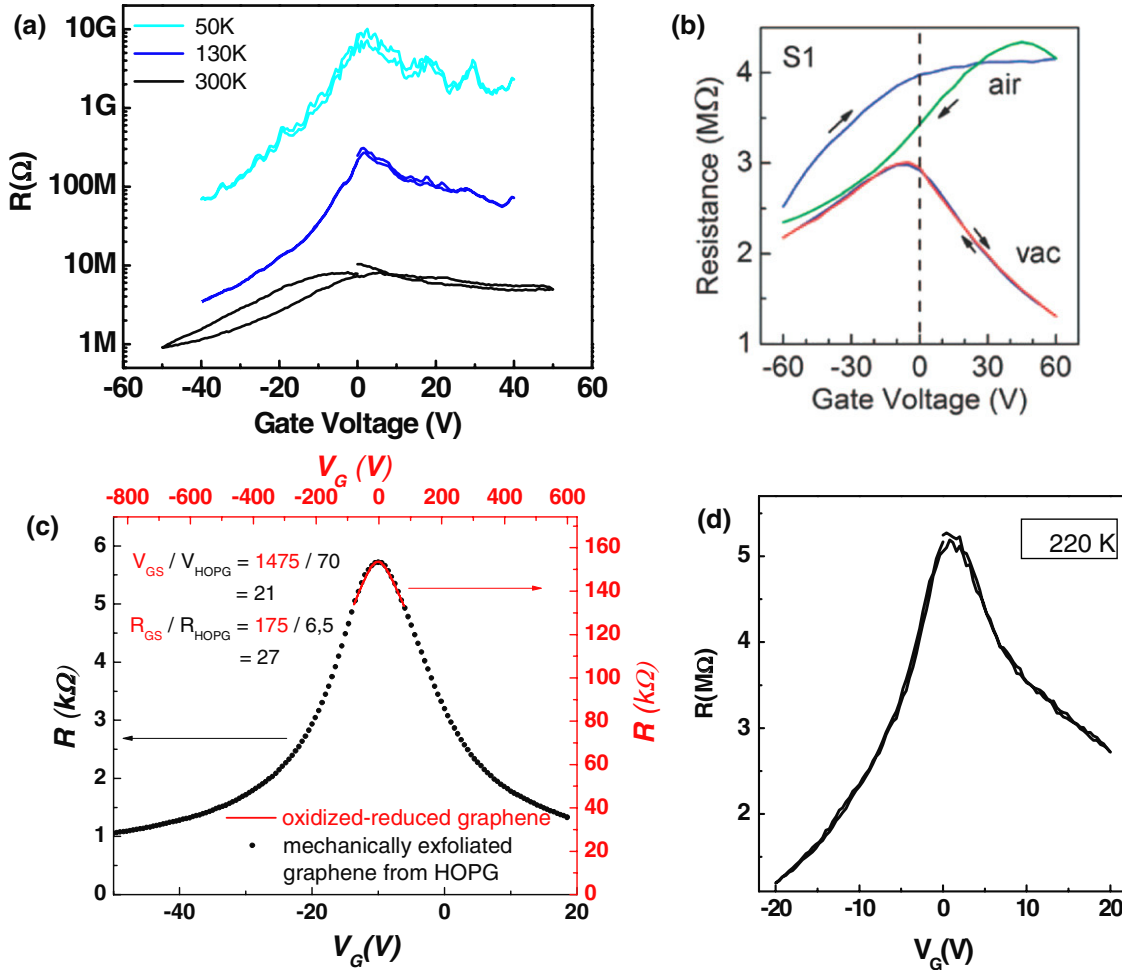


Figure 2. Resistance as a function of back gate voltage of various graphene oxide and RGO monolayers measured in a field effect device configuration (Dirac curve). (a) Reprinted with permission from Gomez-Navarro *et al* [20] Copyright 2007 American Chemical Society. (b) Reprinted with permission from Jung *et al* [27] Copyright 2008 American Chemical Society. (c) Graph reprinted with permission from Gengler *et al* [30] Copyright 2010 Wiley-VCH. (d) Reprinted with permission from Kaiser *et al* [31] Copyright 2009 American Chemical Society.

shift of the maximum towards positive gate voltages, which could be reversed by placing the sample in a vacuum. This observed shift was attributed to doping by oxygen and/or water absorption, similarly to what was observed in the past for carbon nanotubes [26] and for micromechanically cleaved graphene [1]. Similarly, Jung *et al* [27, 28] demonstrated the high sensitivity of the GO to water vapour by recording Dirac curves at various stages of exposure. As shown in figure 2(b), when sweeping the gate voltage there is a large hysteresis effect for the maximum of the Dirac curve of the sample exposed to air. In analogy with water adsorbed on carbon nanotubes [29] this effect has been considered [28] to be due to charge trapping resulting from water (and various microsolvated contaminants) adsorbed both on the graphene oxide sheet and the underlying substrate. Additional treatment in vacuum causes this hysteresis to disappear [27] and drastically alters the electrical properties: the FWHM of the Dirac curve, which is directly connected to the mobility [1], becomes smaller, while the maximum decreases and moves to negative gate voltage, revealing a slight p doping (see figure 2(b)).

Similar observations have been reported for differently functionalized GO films [17, 30–32]. Particularly interesting

are the results on GO layers deposited using a Langmuir–Schaefer method and measured at various stages of treatment, starting from a chemical reduction followed by annealing and exposure of the hot sample to ethylene (in UHV) (see figure 2(c)) [30], where it was shown that this last step triggers not only a further reconstruction of the graphene oxide to graphene but also protects the reduced GO sheets from the environment. In fact, subsequent exposure to air barely affected the electronic properties.

In the quest for best quality chemically prepared graphene, a very recent study by Kaiser *et al* [31], where the GO flakes were chemically reduced in a hydrogen plasma shows the Dirac curve with the smallest FWHM ever observed for reduced GO (figure 2(b)) and with a maximum at zero gate voltage (measurement performed at 220 K). Even this best Dirac curve does not display a clear transition to the pure metallic regime where one should observe a flattening of the curve at high positive or negative gate voltage value since the resistance becomes independent of the applied gate voltage. The best criterion for the quality of reduced GO is the charge carrier mobility μ given by the formula $\mu = (neR)^{-1}$, where n is the carrier density, e is the electron charge and R is the sheet

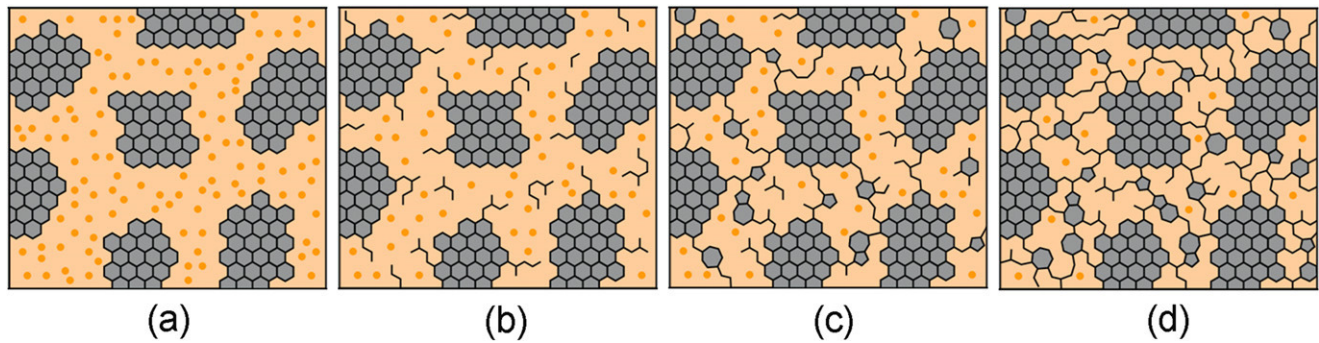


Figure 3. Structural model of graphene oxide at different stages of reduction by thermal annealing taken reprinted with permission from Mattevi *et al* [17] Copyright 2009 Wiley-VCH. (a) Room temperature; (b) $\sim 100^\circ\text{C}$ (c) $\sim 220^\circ\text{C}$; (d) $\sim 500^\circ\text{C}$. The dark grey areas represent sp^2 carbon clusters and the light grey areas represent sp^3 carbon bonded to oxygen groups (represented by small dots). At $\sim 220^\circ\text{C}$, the percolation among the sp^2 clusters initiates (corresponding to sp^2 fraction of ~ 0.6).

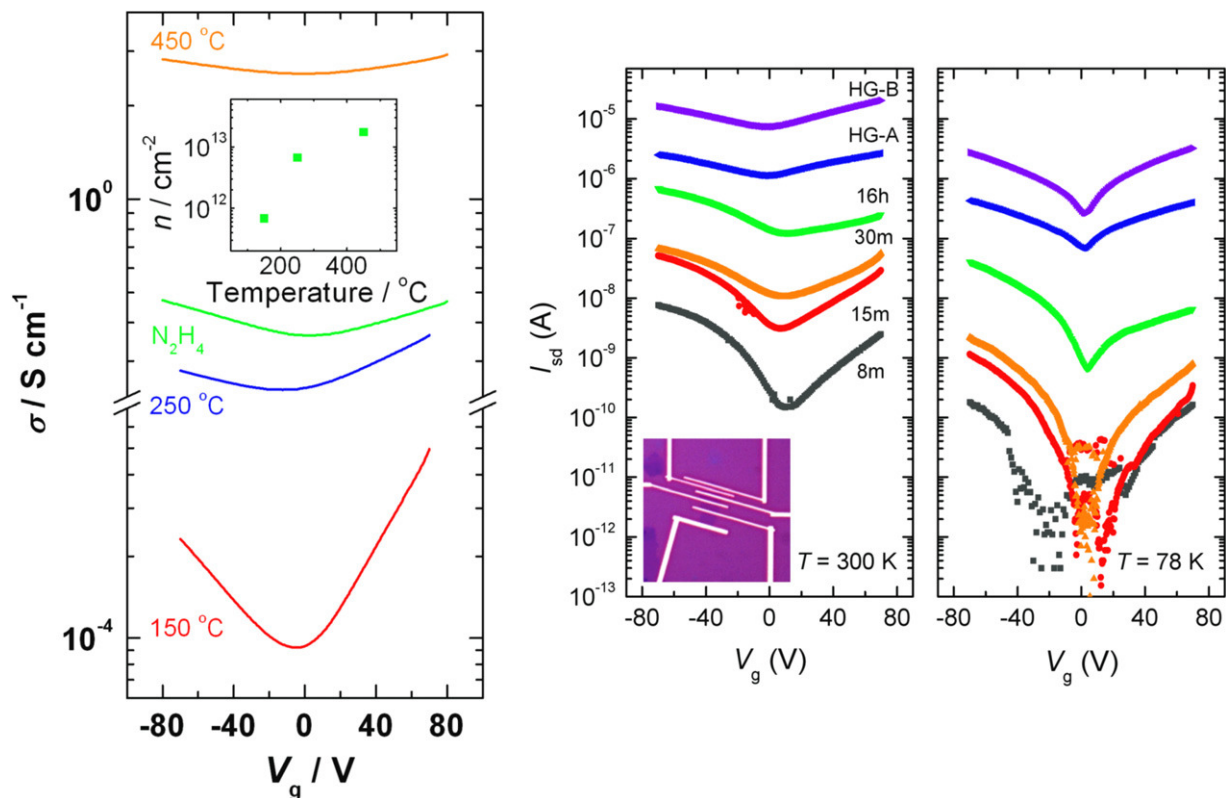


Figure 4. Conductivity (left panel graph reprinted with permission from Mattevi *et al* [17] Copyright 2009 Wiley-VCH) and transfer characteristics (right panel graph reprinted with permission from Eda *et al* [34] Copyright 2009 American Chemical Society) versus gate voltage in field effect devices prepared with graphene oxide as a function of reduction treatment.

resistivity. However, one has to keep in mind that this formula is valid only in the metallic regime where the carrier density can be considered uniform and the presence of electron–hole puddles does not play a role [33]. Using it in the vicinity of the Dirac neutrality point can be misleading. Due to the very broad resistivity curves that are measured in graphene prepared by reducing GO (see figure 1), the applied gate voltages prove not to be enough to reach the metallic regime. To extract the carrier mobility, one can assume that the resistivity versus gate voltage curve resembles the typical resistivity behaviour of micromechanically cleaved graphene in the metallic regime as well, but with a scaling factor for both the gate voltage and the resistivity. As shown in figure 2(c) [30] the comparison

with the measurements on mechanically exfoliated graphene samples and using this scaling yields a carrier mobility for the studied reduced GO in the range of $4\text{ cm}^2\text{ V}^{-1}\text{ s}^{-1}$ in the metallic regime, i.e. a factor of 500 lower than in mechanically exfoliated graphene. From this and from the data presented in figure 2 for other reduced GO, it seems quite clear that the transport properties of reduced GO are limited by structural defects or imperfections. A structural model that captures the essential features of transport through an individual GO sheet at different stages of reduction by thermal annealing was proposed by Mattevi *et al* [17] and is presented as a series of sketches in figure 3. First (figure 3(a)) sp^2 clusters are isolated by areas functionalized with oxygen atoms. As

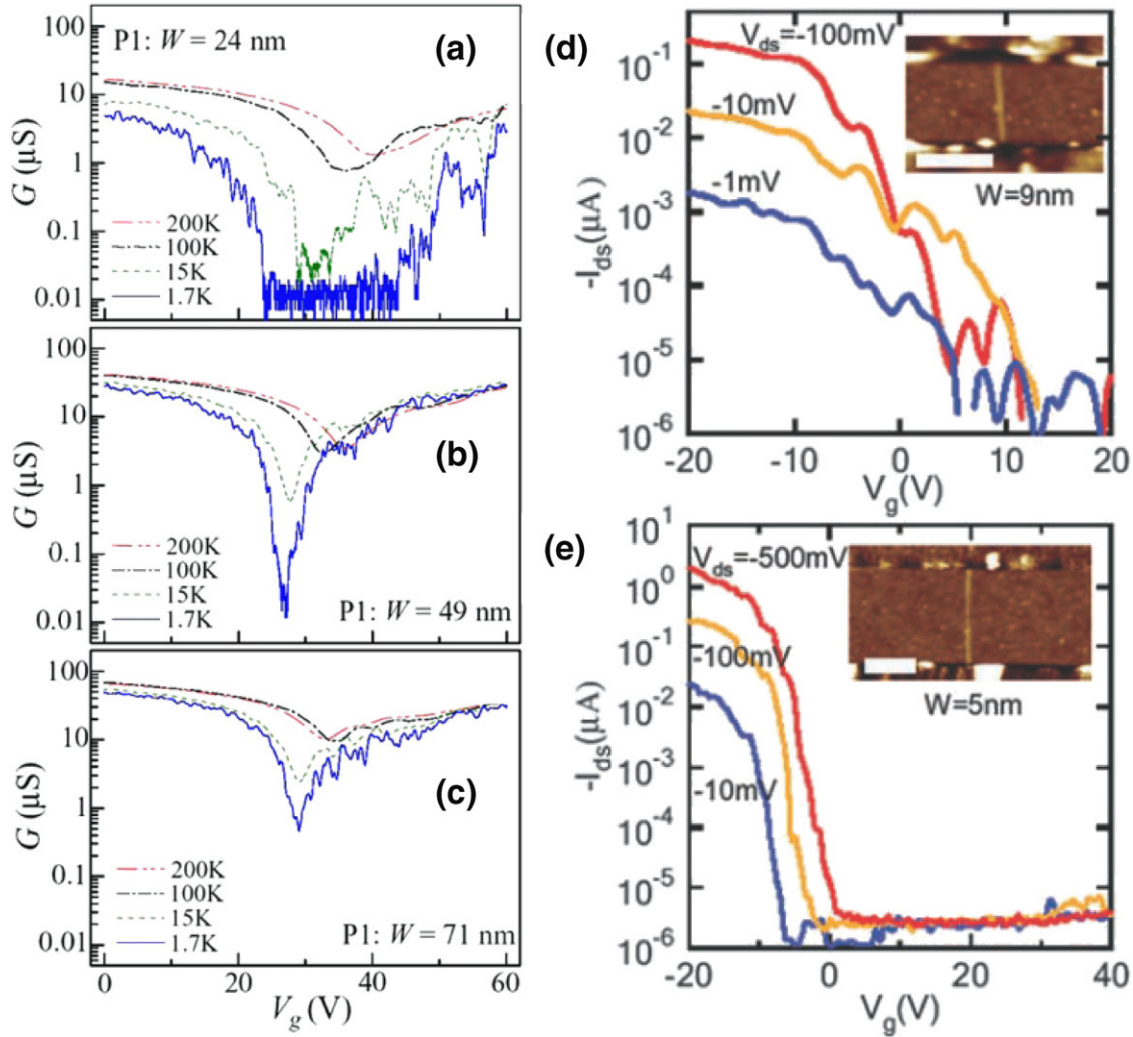


Figure 5. (left panel (a)–(c)) Conductance as a function of gate voltage measured at different temperatures for FETs made from mechanically cleaved graphene which was lithographically patterned into a ribbon structure; figure reprinted with permission from Han *et al* [36] Copyright 2007 American Physical Society. The width of each nanoribbon is indicated in each panel. right panel: transfer characteristics (current versus gate voltage $I_{ds}-V_{gs}$) for FETs built from (d) a 9 nm wide chemically produced nanoribbon (thickness ~ 1.5 nm, \sim two layers) with a channel length $L \sim 130$ nm and (e) transfer characteristics for $a \approx 5$ nm wide nanoribbon (thickness ~ 1.5 nm, \sim two layers and channel length $L \sim 210$ nm) with Pd contacts. The inset shows a AFM image of this device (scale bar is 100 nm). Figure from Li *et al* [35]. Reprinted with permission from AAAS.

the material is progressively reduced, hopping and tunnelling among the clusters increase (figure 3(b)). Further reduction by removal of oxygen leads to greater connectivity among the original graphitic domains by formation of new smaller sp^2 clusters, but also to an increase in structural defects via evolution of CO and/or CO_2 species (especially from epoxy groups), indicated as pentagons in figures 3(c) and (d). Thus, transport in the initial stages of reduction occurs via tunnelling or hopping among the sp^2 clusters. In the following section we shall review what is known about the nature of defects studied by scanning tunnelling microscopy (STM) and high-resolution transmission electron microscopy (HR TEM) but first we would like to focus on how the electronic properties change as a function of the reduction treatment.

An example, shown in the right panel of figure 4, is the work by Eda *et al* [34] who built field effect devices with GO prepared by a modified Hummers and Offeman's method and then reduced by exposure to saturated vapour of hydrazine

monohydrate in a loosely sealed Petri dish at 80°C over a variable length of time. Measurements performed at 78 K and RT (figure 4 right panel) clearly show that the conductivity continuously improves as a function of reduction time. Several GO flakes reduced directly in anhydrous hydrazine prior to deposition were also studied (named HG-A and HG-B in figure 4 (right panel)). The devices made with the latter were annealed in N_2/H_2 (90/10) atmosphere at 150°C for 1 h prior to measurements and exhibit an even better conductivity. However, one also notes that the 'V' shape of the ambipolar graphene transfer characteristics becomes less pronounced as the reduction time progresses. In micromechanically cleaved graphene, this dip in conductivity is well understood and the minimum corresponds to a conductivity $\sim 4e^2/h$ at the charge neutrality point, $V_g = V_{\text{Dirac}}$, where e and h are the electric charge and Planck constant, respectively. A way to quantify the depression of the conductivity dip in reduced GO is to establish the so-called on/off ratio, which is the ratio between the current

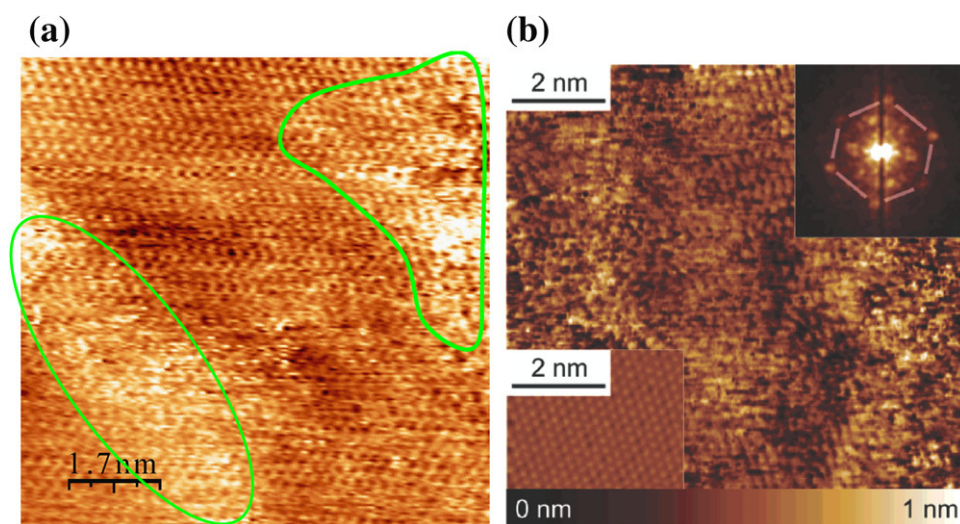


Figure 6. (a) STM image of a graphene oxide monolayer on HOPG. Oxidized regions are marked by green contours, reprinted with permission from Gomez-Navarro *et al* [20] Copyright 2007 American Chemical Society. (b) 10×10 nm STM image of a different type of oxidized graphene on HOPG, Fourier transform of the image shows that the hexagonal order is present (inset on the right top), and inset on the left bottom shows an STM image of HOPG recorded with identical scan conditions, reprinted with permission from Kudin *et al* [21] Copyright 2008 American Chemical Society.

in the ‘supposedly metallic’ regime and the current at the neutrality point. In the data plotted in the right panel of figure 3 a change in the on/off ratio from 10^3 to 2 is observed. Similar behaviour was observed for a three layer GO flake studied by Mattevi *et al* [17] as shown in the left panel of figure 3: as the flake was heated in vacuum to induce reduction thermally, the conductivity went from 10^{-5} to 10 S cm^{-1} but again the ‘V’ shape of the ambipolar graphene transfer characteristics was more pronounced for GO reduced at 150°C compared with films reduced at 450°C and the on/off ratio of the film annealed at 150°C is 10 and decreased to ~ 2 with further reduction. GO reduced by exposure to hydrazine monohydrate vapour at 80°C also exhibited the ‘V’ shape, comparable to devices made with the film annealed at 250°C . These results demonstrate that while higher temperature or longer exposure time to a reducing agent clearly lead to an increased conductivity of the reduced GO sheets, a mild reduction (well tuned timing of exposure to the reducing agent or annealing to around $150\text{--}200^\circ\text{C}$) seems advantageous in order to preserve the ‘V’ shape of the ambipolar graphene transfer characteristics (i.e. a high on/off ratio). Alternatively, improvement can be achieved with the use of ethylene and temperature ranging from 500 to 600°C [30]. However, from all these measurements by various research groups, it seems quite clear that the intrinsic properties of graphene are never completely recovered.

To close this section devoted to the electronic properties of graphene, another proof of the versatility and quality of the chemical approach was given by the impressive results of Li *et al* [35] who deposited graphene nanoribbons with a width of $\sim 10 \text{ nm}$ from a stabilized solution using non-covalent polymer functionalization and integrated them in field-effect transistors to test their low dimensional (quasi-1D) properties. In particular, these nanoribbons were synthesized from commercial expandable graphite (a graphite intercalated with sulfuric acid and nitric acid and hence very similar to the oxidized graphite even if not named so) through exfoliation by

brief (60 s) heating to 1000°C in a mixture of 3% hydrogen in argon. The resulting exfoliated graphite was dispersed in a 1,2-dichloroethane solution of poly(*m*-phenylenevinylene-co-2,5-dioctoxy-*p*-phenylenevinylene) (PmPV) by sonication to form a homogeneous suspension. AFM images of the nanoribbons indicate a height of $1\text{--}1.8 \text{ nm}$ which results from not perfect exfoliation yielding a few layers of graphite and from the non-covalent functionalization with the polymer. In figure 5 we present the transfer characteristics (Dirac curves) of these nanoribbons (right panel) together with those recorded for devices fabricated with mechanically cleaved graphene which was lithographically patterned into a ribbon structure [36]. The left part of figure 5 shows electrical measurements of mechanically cleaved etched graphene [36] resulting in a quasi-one-dimensional structure with narrow width and atomically smooth edges, which exhibits the opening of a bandgap. For such a structure, lateral confinement of charge carriers manifests itself [36] at room temperature with the typical V-shaped conductance versus gate voltage, but showing a minimum conductance of the order of $4e^2/h(W/L)$. However, unlike large graphene sheets, nanoribbons with width $W < 100 \text{ nm}$ show a decrease in minimum conductance of more than one order of magnitude at low temperatures. For example, for the nanoribbon with $W \sim 24 \pm 4 \text{ nm}$ (figure 5(a), left panel), a large ‘gap’ region appears for $25 < V_g < 45 \text{ V}$, where the conductance is below detection limits. This strong temperature dependence differs strongly from that of the large micromechanically cleaved graphene samples where the minimum conductance changes less than 30% in the temperature range $30 \text{ mK--}300 \text{ K}$ [37]. The suppression of the usual V shape curve characteristics of the graphene and its replacement by a clear on/off behaviour around the neutrality point is generated by the opening of the bandgap. Similar transfer characteristics were also observed for the chemically produced graphene nanoribbons (figure 5 right panel) for which Li *et al* [35] observed that the

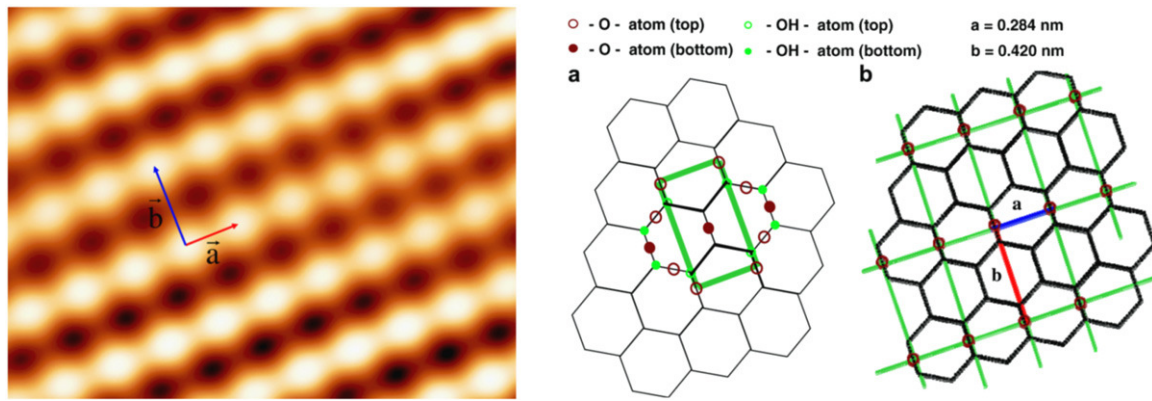


Figure 7. (left panel) STM image, reprinted from Pandey *et al* [40] with permission from Elsevier and model for –O and –OH attachment by Buchsteiner *et al* [39] (right panel). Reprinted from Pandey *et al* [40] with permission from Elsevier.

room-temperature on-off current switching ($I_{\text{on}}/I_{\text{off}}$) induced by the gate voltage increased exponentially with decreasing nanoribbon width. All their measurements gave evidence for p-type unipolar graphene nanoribbon FETs (it is not explained why the behaviour is unipolar) and electrostatic simulations of the gate capacitances gave an estimated hole mobility in the ≤ 10 nm wide ribbons of ~ 100 to $200 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$.

3. Structural integrity of chemically prepared graphene

As shown in the previous section, RGO shows the typical ambipolar behaviour expected for graphene but strongly influenced by the presence of defects/impurities. The structural integrity of the RGO can be investigated by several methods as described hereafter. Low energy electron diffraction and selected area electron diffraction (SAED) have been the methods of choice to reveal local and global high crystallinity of reduced GO and exfoliated samples (see last section for the latter) because the cross section for interaction with electrons is 10^5 higher than that with x-rays, electrons can be focused on very small spots and these techniques do not require access to large-scale facilities like synchrotron or neutron sources. STM and HR TEM are probably today's most powerful tools to investigate, locally, the structural integrity of a given material. Therefore, we review hereafter some of the most relevant studies realized on GO and reduced GO. Figure 6(a) shows the STM image of a GO sheet deposited onto HOPG after dispersion in water with the help of soft sonication [20]. The hexagonal lattice of graphene is clearly preserved in some parts of the sheet while the regions marked by green contours presumably contain functional groups from the oxidation process and appear disordered. The degree of oxidation/defect level of this type of sheets was calculated from a range of measurements and agrees with spectroscopic data [20]. The STM image in figure 6(b) shows similar features for different GOs, called functionalized graphene in the original report, prepared through rapid heating of GO in solution [21] and subsequent deposition on HOPG and is also clearly distinguishable from pristine graphene (shown in the inset of figure 6(b)). Even though figure 6(b) does not reveal large regions with a honeycomb lattice as seen

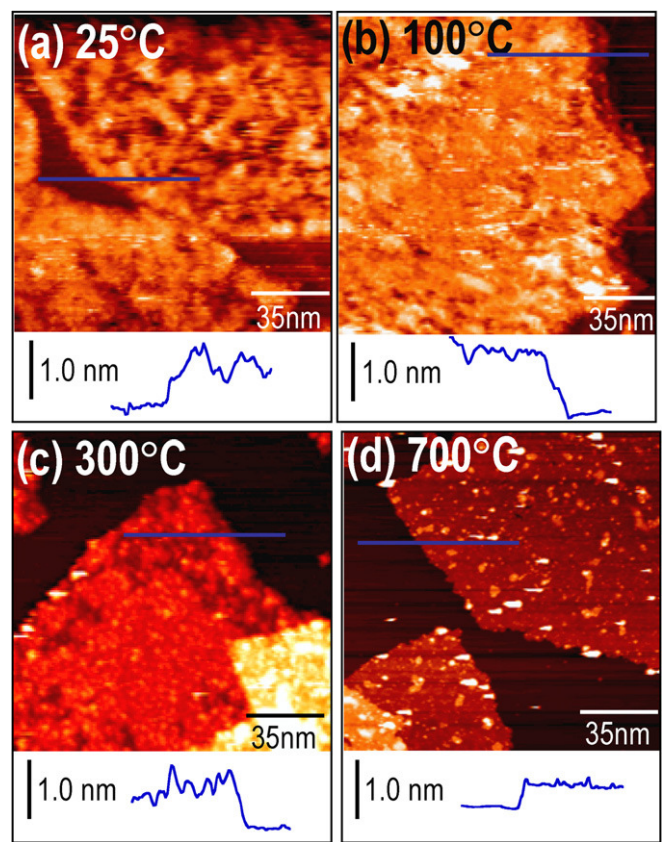


Figure 8. (left) STM image from Chua *et al* [41] of octadecylamine functionalized GO and representative line profile after thermal treatment at different temperatures. (a) Pristine, (b) after 10 min 100°C in N_2 , (c) after 10 min 300°C in N_2 , and (d) after 15 min 700°C in N_2 . Conditions: $V_{\text{tip}} = 2.0 \text{ V}$, $i_{\text{tunnel}} = 100 \text{ pA}$ and $T = 77 \text{ K}$. Image vertical scale: 1.0 nm (bright is high and dark is low). Reprinted with permission from Chua *et al* [41], copyright 2008 American Institute of Physics.

in figure 6(a), a Fourier transform of the image (see inset figure 6(b)) shows a clear hexagonal structure, signature of a graphitic backbone. The observed separation of pristine graphene and disordered regions in these GO sheets agrees with the theoretical prediction that these functional groups should arrange in islands and rows [38]. GO shown in figure 6 appears

quite rough with a peak to peak distance of about 1 nm, most likely due to the presence of *random* –O and –OH or defects. However, a model for the possible bonding sites of –O and –OH on a graphene layer [39] reproduced in figure 7 (right panel) shows a possible arrangement where top and bottom –O and –OH groups are attached to the graphene sheet in a *periodic* fashion.

Such a structure has been observed by Pandey *et al* [40] in STM images of GO prepared following the classic Hummers and Offeman's method as illustrated in the left panel of figure 7. This is the first atomically resolved image of a graphene oxide sheet which seems, nevertheless, to refer to rather small areas since the reported STM field of view is only of $\sim 2 \text{ nm}^2$. This periodic structure can therefore be regarded as a minority phase of GO while the randomly attached functional groups discussed before constitute the majority phase.

Surface functionalization of GO can be a means to introduce desired chemical groups for control of the surface properties and for integration into devices. In this context it is not only important to understand the functionalization process itself but also to identify how the introduced groups influence the graphene properties. As a representative example

[41], STM images of GO nanosheets functionalized with octadecylamine are displayed in figure 8; the line profile across the sheet edge extracted from these images is also shown. The authors argue that by adding the van der Waals radius of graphene to twice that of octadecylamine since the molecules are grafted on both sides of the nanosheet, one obtains a total van der Waals radius of 1.0–1.2 nm, in agreement with the observed STM image. Note that this value also agrees with atomic force microscopy (AFM) data [30] of octadecylamine-functionalized GO. The series of STM images in figure 8 displays the thickness and morphology evolution as a function of the temperature. No changes in morphology/coverage of grafted groups are observed for annealing at 373 K, while annealing beyond this temperature results in a progressive diminution of the bright features on top of graphene, identified as the grafted groups as well as a decrease in the average thickness. This trend is observed up to a temperature of 973 K where very few bright features remain. One can conclude from these observations that the octadecylamine functionalities progressively detach from the graphene sheet.

Additional proof of the graphene-like structure of the graphene oxide and functionalized graphene oxide is provided by TEM images and diffraction patterns (SAED). Figure 9 shows a TEM image of functionalized graphene oxide obtained by reducing exfoliated graphite oxide in the presence of poly(sodium-4 styrene sulfonate) [42]. The large single sheet appears transparent and folded over at the edges, with isolated small graphene fragments on its surface, similarly to what one observes for micromechanically cleaved graphene [43]. Figure 10 presents instead the TEM images and diffraction pattern of graphene oxidized with benzoyl peroxide after reduction [44]. Again single sheets with a hexagonal diffraction pattern are evident. This is the only GO not produced using Hummers and Offeman's, Brodie's or Straudemair's methods and the flakes are significantly smaller than what one can obtain using the 'classical oxidation routes'.

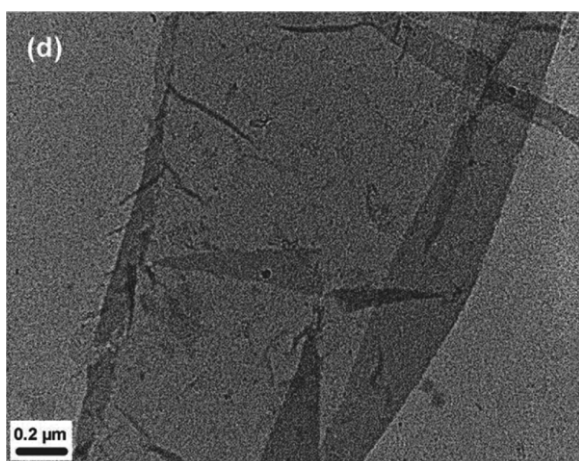


Figure 9. Transmission electron micrograph of water soluble functionalized GO, Reprinted with permission from Si *et al* [42] Copyright 2008 American Chemical Society.

4. Going large scale

While in the previous section we mainly focused on the properties and quality of the graphene flakes prepared by

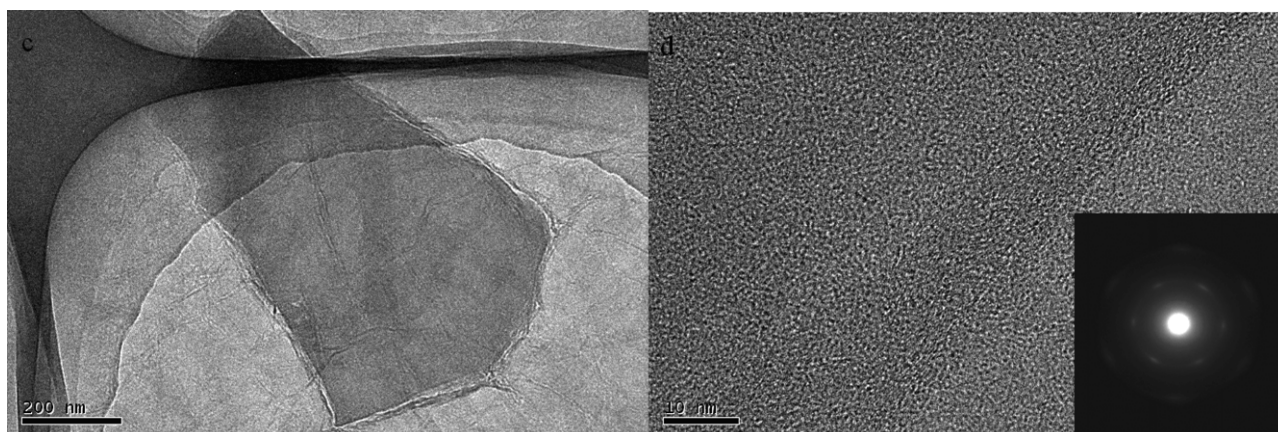


Figure 10. Transmission electron micrographs of benzoyl peroxide oxidized graphene oxide. Scale bar 200 nm (left) and 10 nm (right). Reprinted with permission from Shen *et al* [44] Copyright 2009 American Chemical Society.

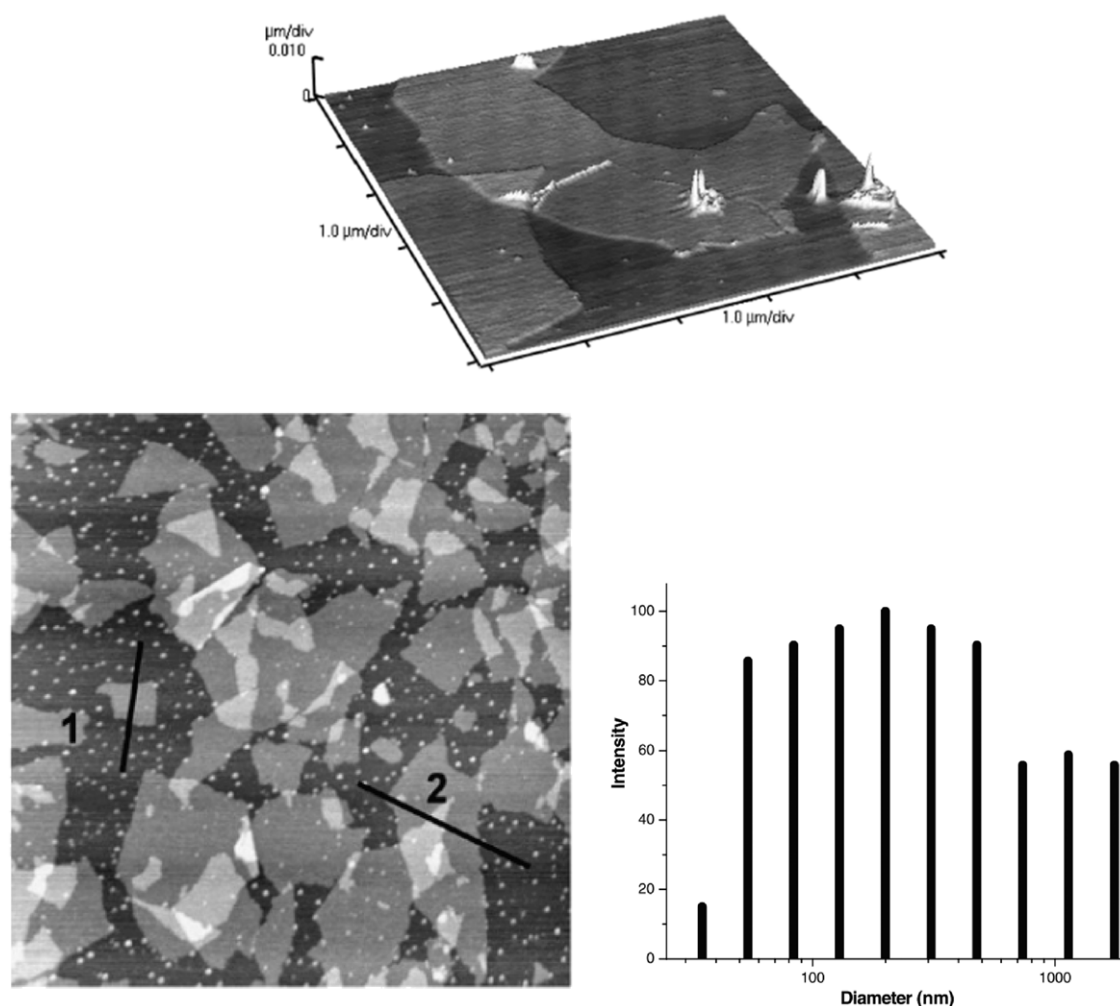


Figure 11. (top) Non-contact mode AFM image of isolated exfoliated single layer graphene oxide; from Stankovich *et al* [45], reproduced with permission from the Royal Society of Chemistry. (bottom) A non-contact AFM image of nanosheets deposited on a mica surface from a dispersion of phenyl isocyanate-treated GO in DMF; line 1 = 0.7 μm , line 2 = 1.0 μm and CONTIN analysis showing the intensity-weighted hydrodynamic diameter distribution of phenyl isocyanate-treated GO nanosheets exfoliated in DMF. Reprinted from Stankovich *et al* [46] with permission from Elsevier.

various methods, in this section we review which approaches can be considered best in terms of ‘large-scale’ production. To this end we compared various preparation methods on the basis of different microscopies or optical images of the produced material. In this comparison we consider first the chemically exfoliated flakes and then how controllably surfaces can be covered with single graphene sheets.

Figure 11 shows one of the first AFM images of isolated exfoliated single layer GO, namely $a \sim 10 \mu\text{m}^2$ sheet deposited from a colloidal suspension onto a mica substrate and measured to be $\sim 1 \text{ nm}$ thick [45]. In the bottom right panel of figure 11, one can see the size distribution of the first chemically derivatized graphite oxide exfoliated in organic solvents (also deposited on mica) [46]. These two examples of single layers of pure and chemically functionalized GO are among the first where complete exfoliation was achieved and have therefore been chosen to illustrate the starting point of a rapid evolution. In these pioneering results neither the coverage nor the number of deposited layers was controlled (see figure 11 top and bottom left). A few years later, still using the same Hummers and Offeman’s method to produce

the starting GO, new deposition procedures were developed to achieve larger flake size, high controllability of the packing and, last but not least, single-layer deposition.

Two examples of optimal results obtained so far for the deposition of chemically prepared graphene are illustrated in figure 12 top and bottom. Cote *et al* [47] used a Langmuir–Blodgett method first reported in Li *et al* [48] but achieved a drastic improvement in terms of flake size and controlled deposition. Similar control alongside with proof of the true graphene quality was reported shortly after by Gengler *et al* [30] (see figure 2). Since one can monitor the surface pressure on top of the water in the Langmuir–Blodgett trough, the packing of the graphene oxide sheets floating at the air/water interface is controlled very easily. As illustrated by the sequence of images (a)–(d) in the bottom panel of figure 12, the more or less compacted floating layer can be transferred to a variety of substrates by vertical or horizontal dipping. One can see an increased coverage; in the first image (figure 12 bottom) where flakes with lateral dimensions of 4–10 μm are well dispersed with a distance of 5–20 μm between them. With increasing surface pressure, the packing increases from

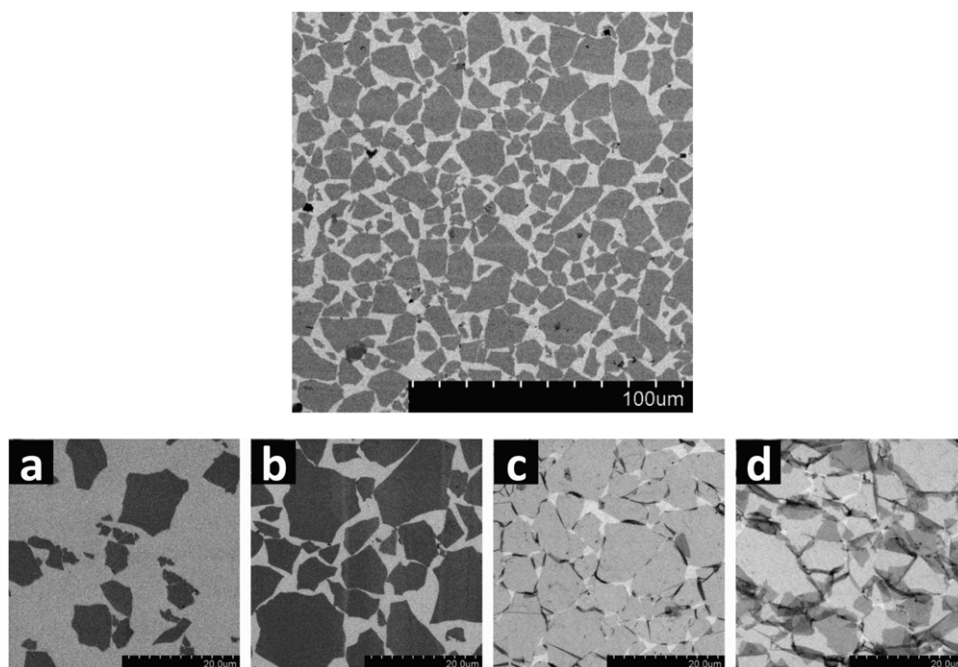


Figure 12. (Top) SEM images of the monolayers of highly covered sample. Scale bars represent 100 μm . (bottom) Langmuir–Blodgett assembly of graphite oxide single layers. (a)–(d) SEM images showing the collected graphite oxide monolayers on a silicon wafer in different regions of the isotherm. The packing density was continuously tuned: (a) dilute monolayer of isolated flat sheets, (b) monolayer of close-packed GO, (c) overpacked monolayer with sheets folded at interconnecting edges and (d) over packed monolayer with folded and partially overlapped sheets interlocking with each other. Scale bars in (a)–(d) represent 20 μm . Image reprinted with permission from Cote *et al* [47] Copyright 2009 American Chemical Society.

image (b), where the inter-flake distance is 1–2 μm to less than 1 μm with some contact between the flakes in image (c), to finally the most dense case in image (d), where the GO flakes are so compressed that they start to overlap at the borders. GO flakes reaching millimetre sizes for use as thin film electrodes when reduced were reported by Su *et al* [49].

A last example of deposition we review here is one where selectivity is achieved through surface modification, namely a study by Wei *et al* [50] of GO transferred from a colloidal dispersion to a patterned substrate. As summarized in figure 13, the authors show that a charged molecular template created using microcontact printing can direct the attachment of a single graphene oxide layer. The method is based on a few simple steps. First the desired template is printed using microcontact printing; the molecule chosen by the authors was 11-amino-1-undecanethiol. Then the gold substrate printed with the pattern of self-assembled molecules is immersed in a GO solution. The authors show that this later step is critical: time, pH and GO concentration influence drastically the quality of the deposition. Once this attachment is achieved, the adsorbed layer can be processed further—for example with a reduction treatment of GO to obtain graphene. Figure 13 documents the success of this method, since in the AFM image on the left (labelled c) selective adsorption can be clearly distinguished. The images on the right of figure 13 show instead the influence of the dipping time on the coverage of the patterned surface, in fact, a dipping time of 5 s (figure 13(a)) produces a coverage of $\sim 50\%$ while immersion for 17 h (figure 13(d)) covered the patterned surface up to $\sim 90\%$. A

similar templated deposition was reported by Li *et al* [51] and in principle one can envision other methods for producing the template such as dip pen lithography [52].

5. The roadmap to high quality—going oxide?

The scientific landscape has been flooded in the few last years by a huge number of reports on new methods of production, deposition, reduction or exfoliation of graphene, each supposedly better than the others. In this section we draw up a condensed summary of the most relevant examples and point out what we believe to be the best currently available ways to prepare graphene. Scheme 1, our roadmap to high quality graphene, displays CVD, mechanical cleavage, Si carbide and other methods defined previously but we shall focus our attention on the methods using chemically derived graphene. Graphite is on top of this scheme and represents today's reality: all or a huge majority of the chemical approaches are top down; in opposition to bottom up approaches so commonly used in material science. Bottom up approaches for graphene-like 2D polymers are not the scope of this review but we refer the reader to Sakamoto *et al* [53].

Anyone getting started in graphene production can choose from many different routes leading to the goal but requiring/involving a series of procedures that will result in the same end material, though of various qualities. Starting from graphite, one has two options, going through oxidation or avoiding it. The easiness and relatively high yield of the graphite oxide approach leads a lot of research groups

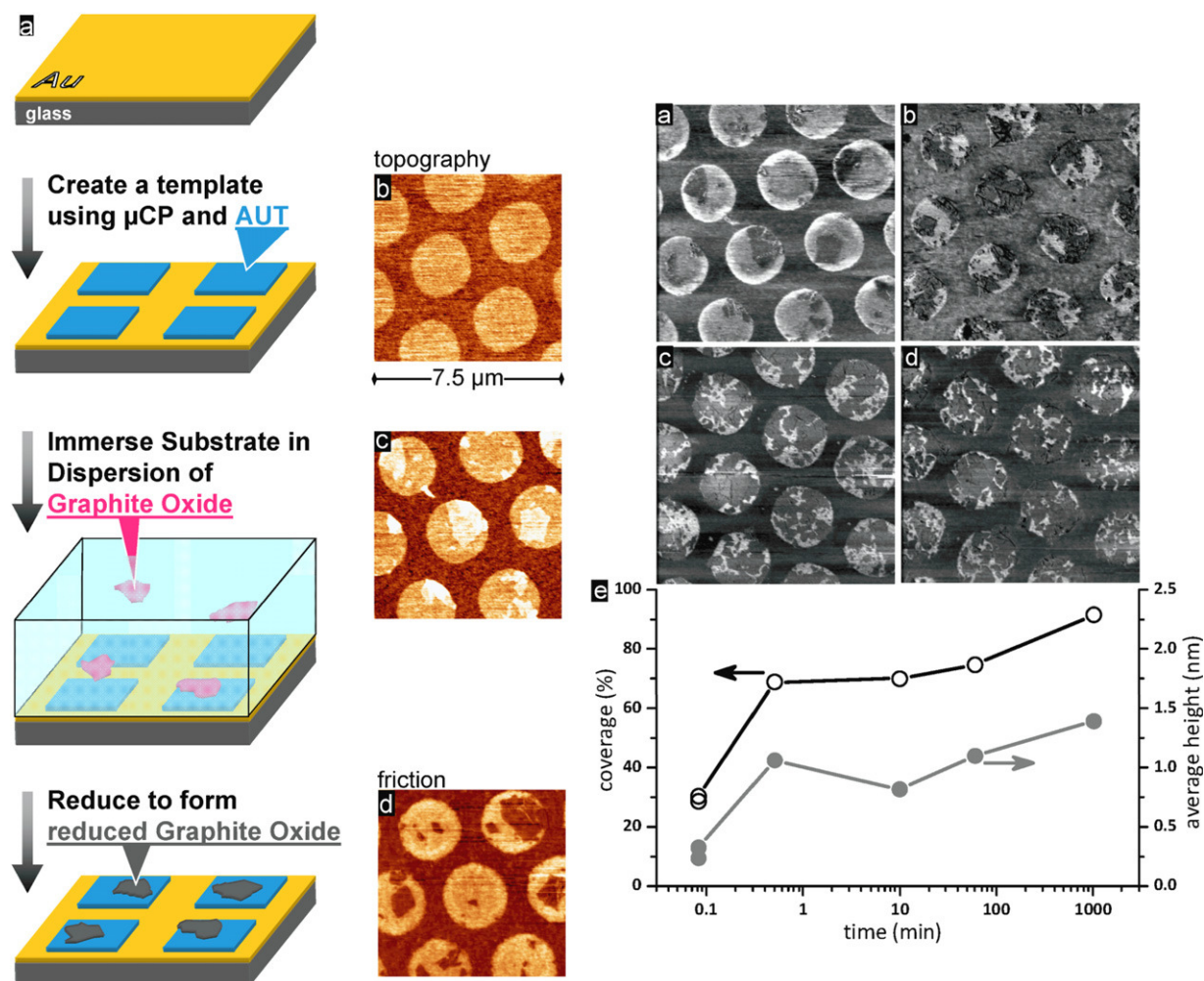
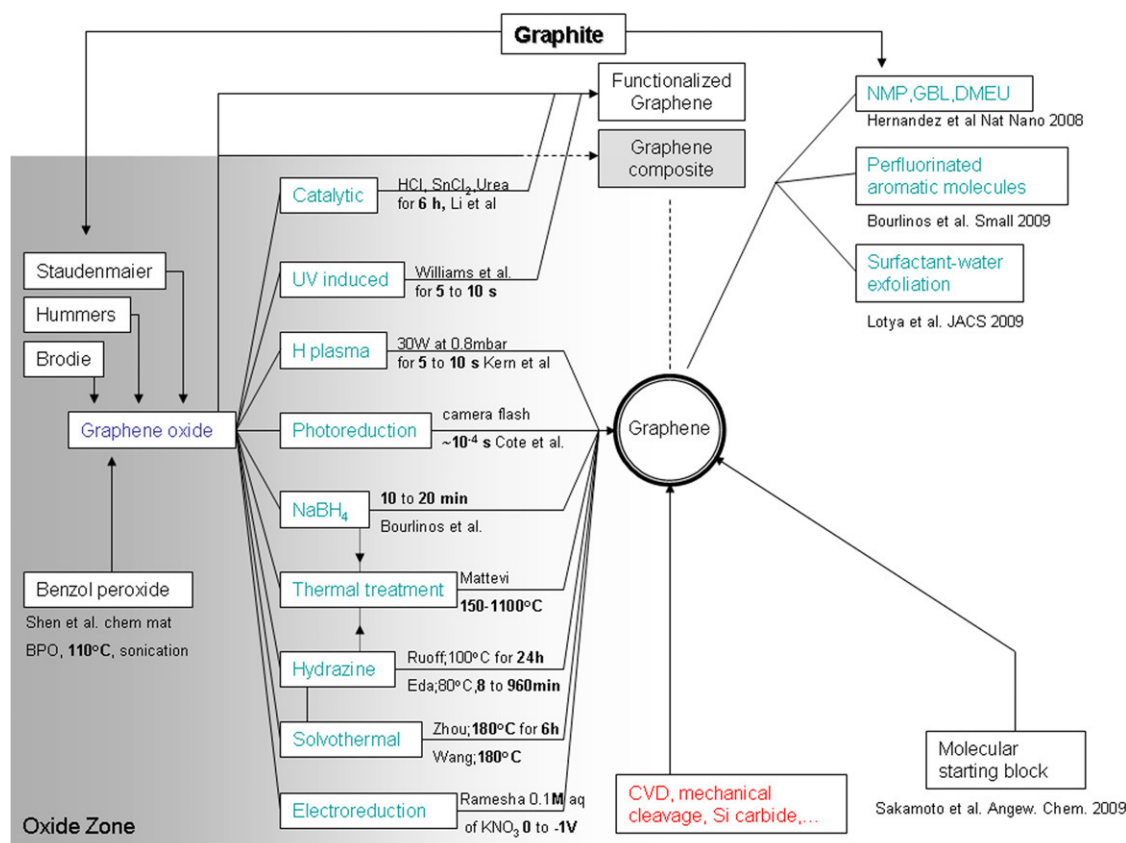


Figure 13. (Left) A scheme of the templating process that shows the formation of the amino-terminated template on mica-peeled gold, followed by immersion in a dispersion of graphite oxide to the reduction of the captured GO to form reduced graphite oxide. (Right (a)–(d)) Friction images of 11-amino 1-undecanethiol (AUT)-patterned Au following 5 s (a), 30 s (b), 10 min (c) and 17 h (d) immersion times in GO dispersions, respectively. All images are $10\ \mu\text{m}$ wide and show the bright (high friction) 11-amino 1-undecanethiol (AUT) being covered with the lower friction GO. (e) Plot of the per cent coverage and average height of the GO films as a function of time. There is a quick adsorption period followed by a much longer and slower adsorption. Reprinted with permission from Wei *et al* [50] Copyright 2008 American Chemical Society.

to follow that path. The starting point in this first case is the oxidation of graphite (any type) by means of potassium chlorate and fuming nitric acid treatment(s) as described by Brodie in 1859 [5] or, as proposed by Staudenmaier [13], by use of sulfuric and nitric acid combined with potassium chlorate or still, as suggested by Hummers and Offeman [14], using concentrated sulfuric acid, sodium nitrate and potassium permanganate as explained in the first section of this review. This last method is the most commonly used for the synthesis of graphite oxide and accepted as the most efficient. To the best of our knowledge, Shen *et al* [44] is the only report of a recent alternative method where benzoyl peroxide is reacted with graphite at moderate temperature and under sonication to produce exfoliated graphene oxide sheets. Depending on the application or fundamental study graphene/graphene oxide is destined for, a number of processing steps is then needed in order to achieve the desired properties, i.e. in many cases to recover a decent conductivity—while still keeping transparency. Other studies report on the creation of composite

material, which either consists of stacked pure GO, GO mixed with a polymer alone or combined with other layered material like clay [54,55]. Some of these composite materials do not require any reduction treatment and use graphene oxide as it is, the GO paper discovered by Ruoff *et al* [56,57] being a famous example of such composite film. On the other hand, the overwhelming majority of the literature of the past few years concentrates on the reconversion of GO to graphene and its deposition for various purposes. Scheme 1 lists the most relevant reduction methods. To the best of our knowledge, the flash photoreduction discovered by Cote *et al* [58] is the fastest reduction process ever (10^{-4} s) and involves, in terms of equipment, only a conventional flash of a digital camera. The reduction occurs because the very large photon flux induces a thermal deoxygenation. Unfortunately, little evidence supports the graphene-like behaviour of the product. Nevertheless, we believe this approach to be promising. Today's most efficient procedure for the reduction of GO seems to be the H plasma reduction method proposed by



Scheme 1. The roadmap to high quality graphene.

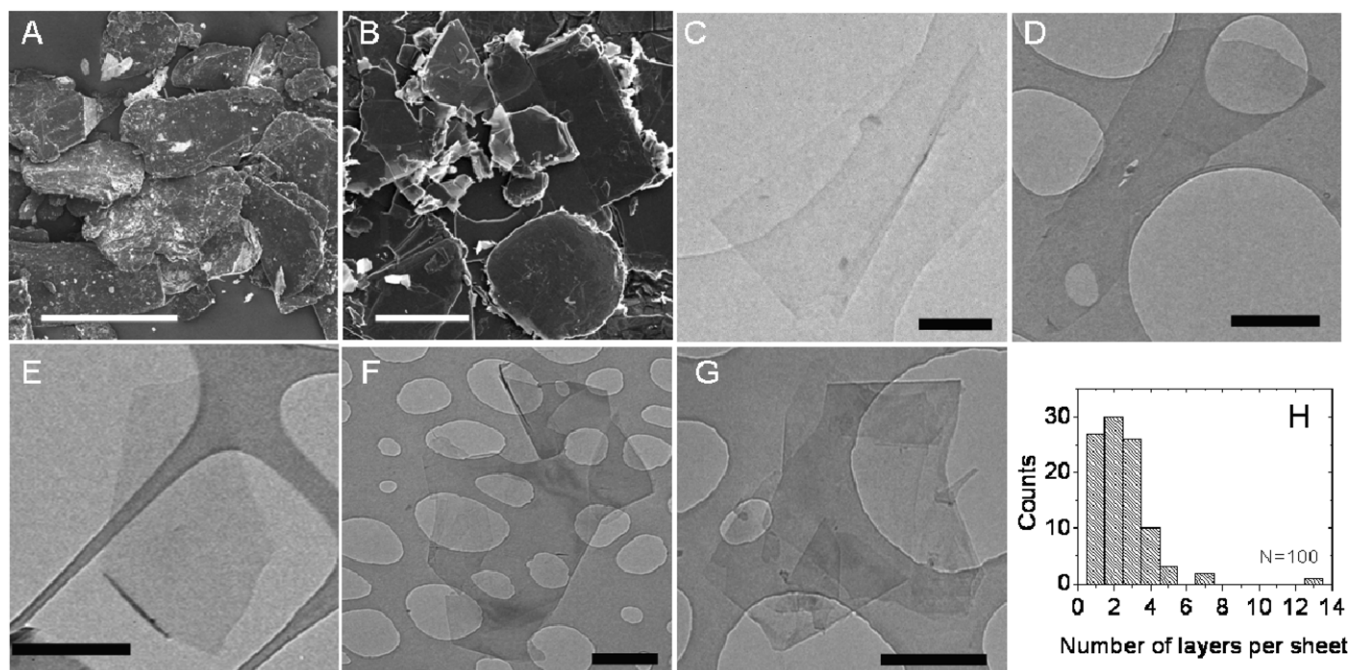


Figure 14. Electron microscopy of graphite and graphene. (a) SEM image of sieved, pristine graphite (scale bar: 500 nm). (b) SEM image of sediment after centrifugation (scale bar: 25 nm). (c)–(e) Bright-field TEM images of monolayer graphene flakes deposited from GBL (c), DMEU (d) and NMP (e), respectively (scale bars: 500 nm). (f), (g) Bright-field TEM images of a folded graphene sheet and multilayer graphene, both deposited from NMP (scale bars: 500 nm). (h) Histogram of the number of visual observations of flakes as a function of the number of monolayers per flake for NMP dispersions. Images and graph reprinted with permission from Macmillan Publishers Ltd: Nature Nanotechnology, Hernandez *et al* [77], Copyright 2008.

Kern *et al* [31], even though it requires more time (5–10 s) and more equipment is needed. This statement is supported by the outstanding Dirac curve displayed in figure 1(b) [31]. The optimized conditions were found to be 5–10 s of exposure to a plasma operating at 0.8 mbar of H and 30 W of power. Ranking next in terms of proven efficiency are purely thermal treatments or heat treatments in vacuum or controlled atmosphere preceded by hydrazine or the NaBH_4 reduction process (for further reading, we recommend recent theoretical works [59, 60] which present possible models for the hydrazine-induced deoxygenation). Once more, the reduction effectiveness is clearly proven by the Dirac curves of devices prepared by various groups ([17, 27, 30, 31, 61–65] see as well figure 1). The most suitable temperature for annealing is still undefined: some groups report on the damage of the ambipolar characteristic after annealing above $\sim 200^\circ\text{C}$ [17], others report on an improved ambipolar behaviour for

temperatures up to $500\text{--}600^\circ\text{C}$ (accompanied by ethylene exposure) [30]. Two additional methods leading to RGO are listed in the lower part of scheme 1: firstly, solvothermal reduction (including hydrothermal reduction) which proceeds in a solvent (water when hydrothermal) while applying mild annealing. The trick is to keep single layers suspended in the solution after reduction. Zhou *et al* [66] showed such an example of hydrothermal reduction and proved, using Raman spectroscopy, x-ray photoelectron spectroscopy (XPS) and AFM that the material was substantially reduced but still single or bilayer. Other examples of solvothermal reduction procedures can be found in the literature [67, 68]; Wang *et al* [69] reported Dirac curves of a solvothermally reduced material, whereas Fan *et al* [70] prepared a graphene suspension through deoxygenation by simply heating an exfoliated-GO suspension under strongly alkaline conditions at moderate temperatures ($50\text{--}90^\circ\text{C}$). Based on these results hydro/solvothermal reduction can be considered a good candidate for reduced GO production if the single layer character can be preserved, for example by depositing the layer beforehand.

An alternative approach reported by Liu *et al* [71], Ganganahalli *et al* [72], Zhou *et al* [73] and Wang *et al* [74] (most probably without knowing of each other's work) is the reduction of graphene oxide using electrochemistry. While voltametry seems to prove an irreversible transformation of the oxidized material, preventing the single layers from aggregating is once more a central issue. Ganganahalli *et al* [72] is the only group actually showing that their electrochemical reduced GO consists of single layers. Although electrochemical methods seem promising, a lot of effort is still needed to develop procedures for a more complete reduction while conserving monolayer characteristics if one actually wants to produce graphene and not very thin graphite!

We complete our review of GO deoxygenation with two examples of catalytic reduction. The first, reported by Williams *et al* [75], describes how graphene oxide suspended in ethanol undergoes reduction as it accepts electrons from UV-irradiated TiO_2 suspensions. The second [76] involves the use of SnCl_2 in HCL and urea, the reaction that creates SnO_2 nanoparticles. The reduced nature of the end product was established by XPS in the SnO_2 case and qualitatively proven by a colour change in the TiO_2 suspensions. The effect of such a nanoparticle overlayer on the electrical properties is unfortunately not documented.

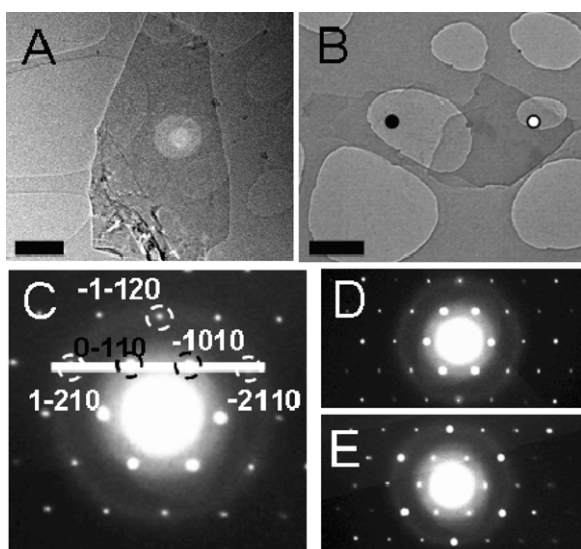
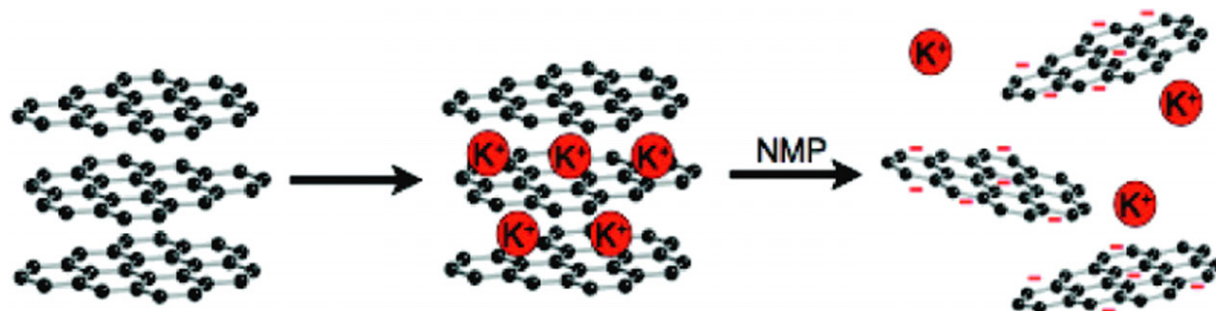


Figure 15. Evidence of monolayer graphene from TEM. (a), (b) High-resolution TEM images of solution-cast monolayer (a) and bilayer (b) graphene (scale bar 500 nm). (c) Electron diffraction pattern of the sheet in (a), with the peaks labelled by Miller-Bravais indices. (d), (e) Electron diffraction patterns taken from the positions of the black (d) and white spots (e), respectively, of the sheet shown in (b), using the same labels as in (c). The graphene is clearly one layer thick in d and two layers thick in (e). Reprinted with permission from Macmillan Publishers Ltd: Nature Nanotechnology, Hernandez *et al* [77], Copyright 2008.



Scheme 2. Representation of the potassium salt $\text{K}(\text{THF}) \times \text{C}_{24}(\text{THF})$ tetrahydrofuran, x (1–3) driven exfoliation of graphite in NMP. Scheme reprinted with permission from Vallés *et al* [80] Copyright 2008 American Chemical Society.

6. Alternative routes

While so far we have concentrated on the ambipolar behaviour, structural properties and the coverage analysis concerning RGO, this section is devoted to the chemical preparation of graphene without oxidation (listed in the right part of our roadmap to graphene (scheme 1)). A very effective method for the exfoliation of graphite with the help of organic solvents such as *N*-methylpyrrolidone (NMP), *N,N*-dimethylacetamide (DMA), *g*-butyrolactone (GBL) and 1,3-dimethyl-2-imidazolidinone (DMEU) was proposed by

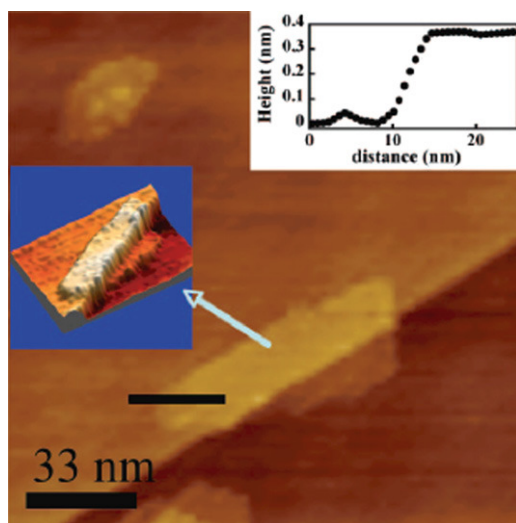


Figure 16. Ambient STM image of a file-down graphite deposit drop casted from solution on a HOPG substrate, showing a graphene flake, lying on a HOPG step. Height scan inset shows a height difference of 0.36 nm between substrate and flake. Similar results have been obtained on a vast number of flakes. Image reprinted with permission from Vallés *et al* [80] Copyright 2008 American Chemical Society.

Hernandez *et al* [77]. This exfoliation takes place because the energy required to exfoliate graphene matches the solvent–graphene interaction energy and extra energy provided through sonication activates the process. It works for solvents which have an interaction energy with graphene that is equal to graphene–graphene interaction energy [1, 78, 79], resulting in a minimal energy cost to overcome the Van Der Waals forces between the graphene sheets. High quality graphene is produced in this way but the very low concentration of single layer graphene in the suspension (0.01 mg ml^{-1}) [77] is a drawback. Some of the best TEM images using the solvent exfoliation route are displayed in figure 14 where one can distinguish monolayer graphene with sizes in the range 0.5–1 μm . Among the proposed solvents NMP seems to yield the best results since one can clearly see graphene single layers in the corresponding TEM images. Figure 14(h) displays a histogram of the number of layers per sheet for the exfoliation of graphite in NMP which is peaked between 1 and 4 layers.

SAED of the exfoliated material is in figure 15, and shows a hexagonal pattern illustrative of the hexagonal carbon honeycomb arrangement of graphite. The existence of monolayer and bilayer graphene arrangements is as well demonstrated by $\{2\ 1\ 1\ 0\}$ spots appearing more intense relative to the $\{1\ 1\ 0\}$.

An alternative route, still NMP based but avoiding the sonication step to afford bigger flakes, was proposed by Vallés *et al* [80]. Inspired by carbon nanotube processing technology [81], they showed that the exposure of graphite to a ternary potassium salt $\text{K}(\text{THF}) \times \text{C}_{24}(\text{THF})$ tetrahydrofuran, $\times (1-3)$ in NMP leads to stable exfoliation of graphite. As depicted in scheme 2, an alkali metal graphite intercalation compound is formed, which, thanks to its charge, helps the exfoliation of graphite in NMP and leads to a stable suspension of negatively charged graphene in NMP mixed with the alkaline salt.

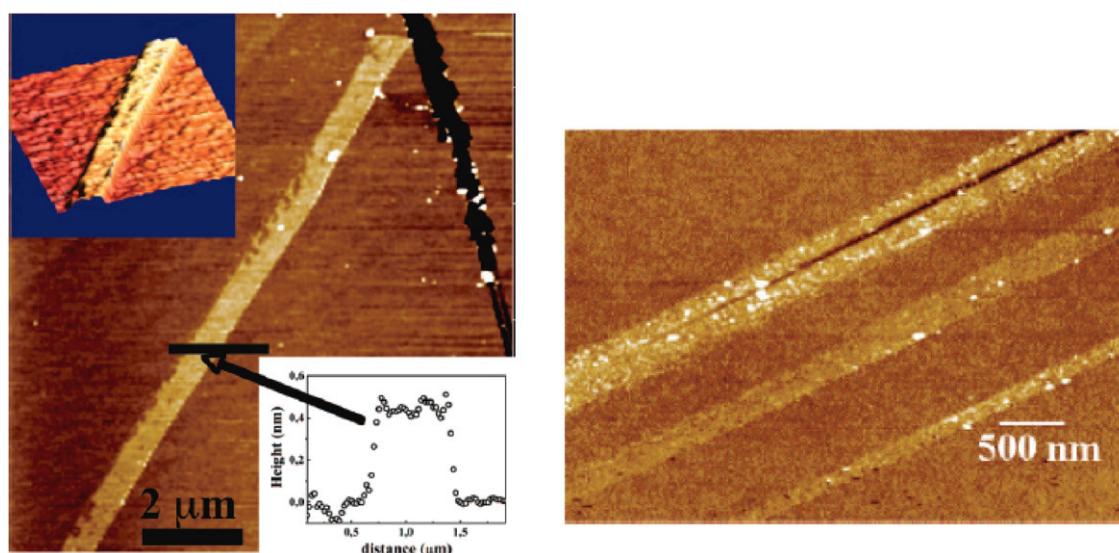


Figure 17. Tapping mode AFM image of a deposit performed by dip-coating of a graphene solution (from expanded graphite) onto mica. Height measurements of the ribbon show a height of 0.4 nm; the full length of the ribbon is about 40 μm . (Right) tapping mode AFM image of a deposit performed by dip-coating of a graphene solution (from expanded graphite) onto Si/SiO_2 wafer. Height measurements give between 0.8 and 1.3 nm for the three ribbons (multilayer). Image reprinted with permission from Vallés *et al* [80] Copyright 2008 American Chemical Society.

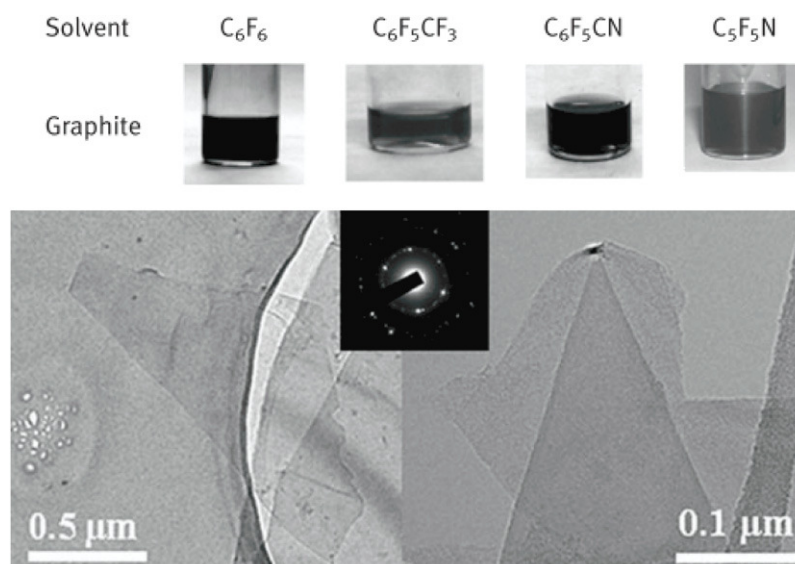


Figure 18. (Top) Colloidal dispersions obtained after liquid-phase exfoliation of graphite using the perfluorinated aromatic solvents below. (Bottom) TEM images of some pentafluorobenzonitrile-etched thin sheets. The SAED pattern is included as inset. Images reprinted with the permission from Bourlinos *et al* [82] Copyright 2009 Wiley-VCH.

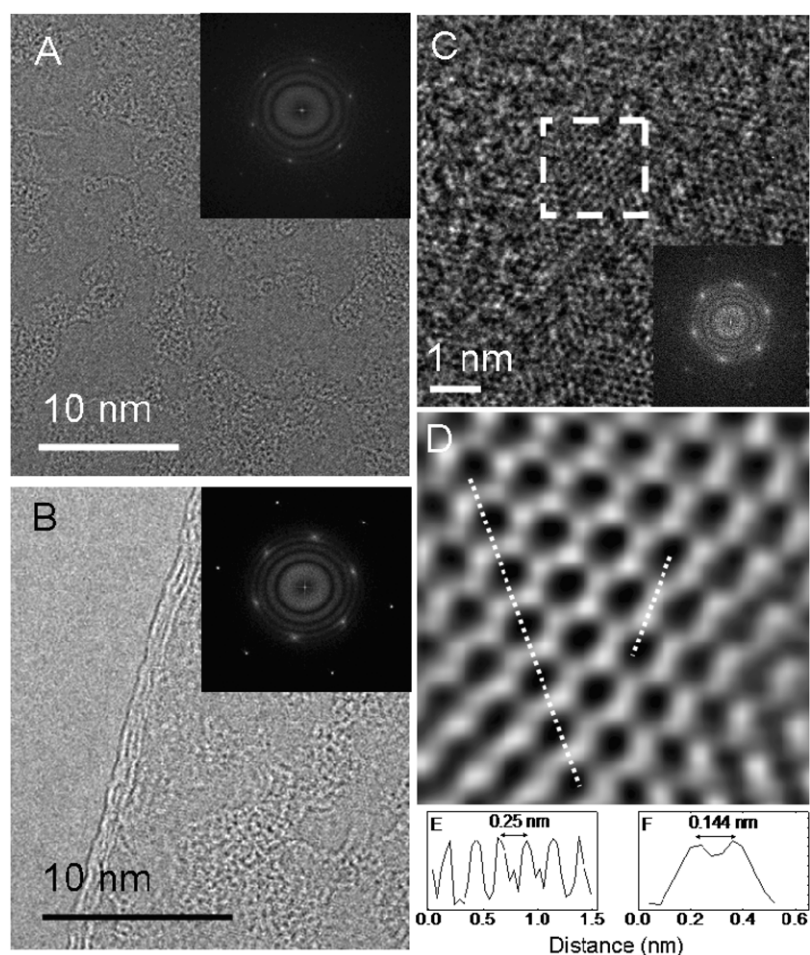


Figure 19. High-resolution TEM images of surfactant exfoliated graphene flakes. (a) A HRTEM image of a section of a graphene monolayer. Inset: fast Fourier transform (equivalent to an electron diffraction pattern) of the image. (b) HRTEM image of a section of a trilayer. Inset: fast Fourier transform of the image. (c) HRTEM image of part of a graphene monolayer. Inset: fast Fourier transform of the region enclosed by the white square. The scale bar is 1 nm. (d) A filtered image of part of the region in the white square. (e) Intensity analysis along the left white dashed line shows a hexagon width of 2.4 Å. (f) Intensity analysis along the right white dashed line shows a C–C bond length of 1.44 Å. Reprinted with permission from Lotya *et al* [85] Copyright 2009 American Chemical Society.

As expected for this mild exfoliation method, yields drastically improved flake size as compared with the results reported in figures 14 and 15. One of the best STM images of chemically exfoliated graphene produced without oxidation is shown in figure 16. The height of 0.3 nm confirms the single-layer character of the graphene flake. Tapping mode AFM images show large-scale monolayer graphene ribbons as reported in figure 17. Again, a height of 0.3 nm was measured on the ribbon. Unfortunately no characterization of the electrical properties was performed on the produced material.

Very recently Bourlinos *et al* [82] presented an approach following the idea of Hernandez *et al* [77] for the exfoliation of graphite with the help of solvents with a surface energy that matches the graphene–graphene interaction energy [1, 78, 79]. Some of the solvents of this approach belong to a peculiar class of perfluorinated aromatic molecules [83, 84] and include hexafluorobenzene (C_6F_6), octafluorotoluene ($C_6F_5CF_3$), pentafluorobenzonitrile (C_6F_5CN) and pentafluoropyridine (C_5F_5N). Aside from the aromatic compounds, Bourlinos *et al* also found that some non-aromatic solvents tested successfully for dispersing graphite, namely ethyl acetate, vinyl acetate, methyl chloroacetate, 2-methoxyethyl ether, acetylacetone and *N,N,N',N'*-tetramethylmethylenediamine, which all exhibited remarkable colloidal stabilities and concentrations of 0.2–0.3 mg ml⁻¹. Examples of the dispersions and TEM images are presented in figure 18. Uniform dark dispersions testify to the absence of aggregation, while single to 2–3 layer thick graphene with flake size in the order of a few micrometres were observed by TEM.

Lotya *et al* [85] proposed a liquid-phase exfoliation of graphene which does not require oxidation or high temperature, avoids expensive solvents and is even, according to the author, safe and user friendly. Here graphite was dispersed in surfactant-water solutions in a manner similar to surfactant-aided carbon nanotube dispersion [86–90]. The dispersed graphitic/graphene flakes are stabilized against reaggregation by Coulomb repulsion between the adsorbed surfactant molecules. Electrical measurements on such flakes show a conductivity of 35 S m⁻¹, a low value attributed to the presence of residual surfactant molecules which are difficult to remove even after several washing treatments. After annealing at 250 °C, as expected, the conductivity rose to 1500 S m⁻¹ (the sheet resistance fell from 920 to 22.5 K Ω) while optical measurements demonstrated that the transparency did not vary throughout the processing. However, this conductivity value is still far from those resulting after the reduction of graphene oxide, which vary from 7200 [91] to 10000 S m⁻¹ [16] and is also significantly lower than the conductivity of graphene derived from exfoliation in NMP (6500 S m⁻¹) [77]. Nevertheless, HR TEM with atomic resolution reproduced in figure 19(a) shows the monolayers to be well graphitized and largely defect free. The authors observed large flakes were of monolayers and bilayers but also reaggregation of thin layers. SAED images reveal the hexagonal lattice of graphene and from the analysis of images like that reproduced in figure 19(b) one can deduce from the number of lines in the edge of the flake that it is a three layer graphite sheet [92]. The

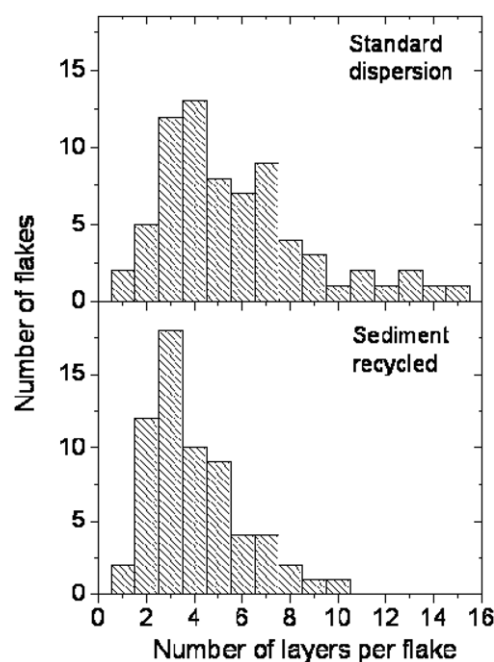


Figure 20. Histograms of the number of layers per flake for dispersions from original sieved graphite and from recycled sediment graphs reprinted with permission from Lotya *et al* [85] Copyright 2009 American Chemical Society.

relative amount of the multilayer structures is illustrated in the histogram of the number of layers per flake for dispersions from original sieved graphite and from recycled sediment shown in figure 20 (the very large flakes are ignored in this histogram) from which one deduces that ~43% of flakes have less than 5 layers and about ~3% of the flakes were monolayer graphene. While this value is considerably smaller than that observed for graphene/solvent dispersions, working in aqueous systems has its own advantages. In general, the majority of these few-layer flakes had lateral dimensions of ~1 μ m. Although these results do not compete with the top quality of the RGO, we believe that today's research is only a beginning and much more can be expected from such an approach. Characterization of the electrical properties would be of great help to learn more about the true quality of the prepared material and it is obvious that efforts are still needed to conserve larger flake sizes with this kind of approach.

7. Conclusion

We reviewed the chemical approaches to graphene production, their advantages as well as their downsides. In doing so, we have drawn a roadmap of today's most reliable path to high-quality graphene via chemical preparation, hopefully helping researchers to find their path through this huge diversity of approaches. This roadmap also reveals the weak points of today's knowledge regarding chemically prepared graphene and points out directions for further research effort. For additional reading, we refer the reader to a review focused on graphene oxide by Compton and Nguyen [93], a general overview by Allen *et al* [94] and one on chemistry and functionalization of graphene by Loh *et al* [95].

References

- [1] Novoselov K S *et al* 2004 Electric field effect in atomically thin carbon films *Science* **22** 666–9
- [2] Lee C, Wei X, Kysar J W and Hone J 2008 Measurement of the elastic properties and intrinsic strength of monolayer graphene *Science* **18** 385–8
- [3] Balandin A A *et al* 2008 Superior thermal conductivity of single-layer graphene *Nano Lett.* **8** 902–7
- [4] Nair R R *et al* 2008 Fine structure constant defines visual transparency of graphene *Science* **6** 1308
- [5] Brodie B C 1859 On the atomic weight of graphite *Phil. Trans. R. Soc. Lond.* **149** 249–9
- [6] Kim K S *et al* 2009 Large-scale pattern growth of graphene films for stretchable transparent electrodes *Nature* **5** 706–10
- [7] Van Bommel A, Crombeen J and Van Tooren A 1975 LEED and Auger electron observations of the SiC(000 1) surface *Surf. Sci.* **2** 463–72
- [8] Forbeaux I, Themlin J and Debever J 1998 Heteroepitaxial graphite on 6H–SiC(000 1): interface formation through conduction-band electronic structure *Phys. Rev. B* **15** 16396
- [9] Berger C *et al* 2004 Ultrathin epitaxial graphite: 2D electron gas properties and a route toward graphene-based nanoelectronics *J. Phys. Chem. B* **1** 19912–16
- [10] Seyller T, Emtsev K V, Speck F, Gao K and Ley L 2006 Schottky barrier between 6H–SiC and graphite: implications for metal/SiC contact formation *Appl. Phys. Lett.* **88** 242103
- [11] Emtsev K V *et al* 2009 Towards wafer-size graphene layers by atmospheric pressure graphitization of silicon carbide *Nature Mater.* **8** 203–7
- [12] Schafhaeuti C 1840 *Phil. Mag.* **16** 570–90
- [13] Staudenmaier L 1898 Verfahren zur Darstellung der Graphitsäure *Ber. Deutsch. Chem. Ges.* **31** 1481–7
- [14] Hummers W S and Offeman R E 1958 Preparation of graphitic oxide *J. Am. Chem. Soc.* **1** 1339
- [15] Cai W *et al* 2008 Synthesis and solid-state NMR structural characterization of ¹³C-labeled graphite oxide *Science* **26** 1815–7
- [16] Becerril H A, Mao J, Liu Z, Stoltenberg R M, Bao Z and Chen Y 2008 Evaluation of solution-processed reduced graphene oxide films as transparent conductors *ACS Nano* **1** 463–70
- [17] Mattevi C *et al* 2009 Evolution of electrical, chemical, and structural properties of transparent and conducting chemically derived graphene thin films *Adv. Funct. Mater.* **19** 2577–83
- [18] Yang D *et al* 2009 Chemical analysis of graphene oxide films after heat and chemical treatments by x-ray photoelectron and micro-Raman spectroscopy *Carbon* **47** 145–52
- [19] Mkhoyan K A *et al* 2009 Atomic and electronic structure of graphene-oxide *Nano Lett.* **9** 1058–63
- [20] Gomez-Navarro C *et al* 2007 Electronic transport properties of individual chemically reduced graphene oxide sheets *Nano Lett.* **7** 3499–503
- [21] Kudin K N, Ozbas B, Schniepp H C, Prud'homme R K, Aksay I A and Car R 2008 Raman spectra of graphite oxide and functionalized graphene sheets *Nano Lett.* **8** 36–41
- [22] Dreyer D R, Park S, Bielawski C W and Ruoff R S 2010 The chemistry of graphene oxide *Chem. Soc. Rev.* **39** 228
- [23] Park S and Ruoff R S 2009 Chemical methods for the production of graphenes *Nature Nanotechnol.* **4** 217–24
- [24] Karaborni S, Smit B, Heidug W, Urai J and Oort E V 1996 The swelling of clays: molecular simulations of the hydration of montmorillonite *Science* **271** 1102–4
- [25] Bolotin K *et al* 2008 Ultrahigh electron mobility in suspended graphene *Solid State Commun.* **146** 351–5
- [26] Martel R *et al* 2001 Ambipolar electrical transport in semiconducting single-wall carbon nanotubes *Phys. Rev. Lett.* **87** 256805
- [27] Jung I, Dikin D A, Piner R D and Ruoff R S 2008 Tunable electrical conductivity of individual graphene oxide sheets reduced at 'low' temperatures *Nano Lett.* **8** 4283–87
- [28] Jung I, Dikin D, Park S, Cai W, Mielke S L and Ruoff R S 2008 Effect of water vapor on electrical properties of individual reduced graphene oxide sheets *J. Phys. Chem. C* **112** 20264–8
- [29] Kim W, Javey A, Vermesh O, Wang Q, Li Y and Dai H 2003 Hysteresis caused by water molecules in carbon nanotube field-effect transistors *Nano Lett.* **3** 193–8
- [30] Gengler R Y N *et al* 2010 Large-yield preparation of high-electronic-quality graphene by a Langmuir–Schaefer approach *Small* **6** 35–9
- [31] Kaiser A B, Gómez-Navarro C, Sundaram R S, Burghard M and Kern K 2009 Electrical conduction mechanism in chemically derived graphene monolayers *Nano Lett.* **9** 1787–92
- [32] Wang S *et al* 2008 Band-like transport in surface-functionalized highly solution-processable graphene nanosheets *Adv. Mater.* **20** 3440–6
- [33] Martin J *et al* 2008 Observation of electron–hole puddles in graphene using a scanning single-electron transistor *Nature Phys.* **4** 144–8
- [34] Eda G, Mattevi C, Yamaguchi H, Kim H and Chhowalla M 2009 Insulator to semimetal transition in graphene oxide *J. Phys. Chem. C* **113** 15768–71
- [35] Li X, Wang X, Zhang L, Lee S and Dai H 2008 Chemically derived, ultrasmooth graphene nanoribbon semiconductors *Science* **319** 1229–32
- [36] Han MY, Ozyilmaz B, Zhang Y and Kim P 2007 Energy band-gap engineering of graphene nanoribbons *Phys. Rev. Lett.* **98** 206805
- [37] Tan Y, Zhang Y, Stormer H L and Kim P 2007 Temperature dependent electron transport in graphene *Eur. Phys. J.—Spec. Top.* **148** 15–18
- [38] Li J, Kudin K N, McAllister M J, Prud'homme R K, Aksay I A and Car R 2006 Oxygen-driven unzipping of graphitic materials *Phys. Rev. Lett.* **96** 176101
- [39] Buchsteiner A, Lerf A and Pieper J 2006 Water dynamics in graphite oxide investigated with neutron scattering *J. Phys. Chem. B* **110** 22328–38
- [40] Pandey D, Reifengerger R and Piner R 2008 Scanning probe microscopy study of exfoliated oxidized graphene sheets *Surf. Sci.* **602** 1607–13
- [41] Chua L *et al* 2008 Deoxidation of graphene oxide nanosheets to extended graphenites by 'unzipping' elimination *J. Chem. Phys.* **129** 114702–6
- [42] Si Y and Samulski E T 2008 Synthesis of water soluble graphene *Nano Lett.* **8** 1679–82
- [43] Meyer J C, Girit C O, Crommie M F and Zettl A 2008 Imaging and dynamics of light atoms and molecules on graphene *Nature* **454** 319–22
- [44] Shen J *et al* 2009 Fast and facile preparation of graphene oxide and reduced graphene oxide nanoplatelets *Chem. Mater.* **21** 3514–20
- [45] Stankovich S, Piner R D, Chen X, Wu N, Nguyen S T and Ruoff R S 2006 Stable aqueous dispersions of graphitic nanoplatelets via the reduction of exfoliated graphite oxide in the presence of poly(sodium 4-styrenesulfonate) *J. Mater. Chem.* **16** 155–8
- [46] Stankovich S, Piner R D, Nguyen S T and Ruoff R S 2006 Synthesis and exfoliation of isocyanate-treated graphene oxide nanoplatelets *Carbon* **44** 3342–47
- [47] Cote L J, Kim F and Huang J 2009 Langmuir–Blodgett assembly of graphite oxide single layers *J. Am. Chem. Soc.* **131** 1043–9

- [48] Li X *et al* 2008 Highly conducting graphene sheets and Langmuir–Blodgett films *Nature Nanotechnol.* **3** 538–42
- [49] Su C *et al* 2009 Electrical and spectroscopic characterizations of ultra-large reduced graphene oxide monolayers *Chem. Mater.* **21** 5674–80
- [50] Wei Z, Barlow D E and Sheehan P E 2008 The assembly of single-layer graphene oxide and graphene using molecular templates *Nano Lett.* **8** 3141–5
- [51] Li B, Lu G, Zhou X, Cao X, Boey F and Zhang H 2009 Controlled assembly of gold nanoparticles and graphene oxide sheets on dip pen nanolithography-generated templates *Langmuir* **25** 10455–8
- [52] Wang Y, MasPOCH D, Zou S, Schatz G C, Smalley R E and Mirkin C A 2006 Controlling the shape, orientation, and linkage of carbon nanotube features with nano affinity templates *Proc. Natl Acad. Sci. USA* **103** 2026–31
- [53] Sakamoto J, vanHeijst J, Lukin O and Schlüter A 2009 Two-dimensional polymers: just a dream of synthetic chemists? *Angew. Chem. Int. Edn Engl.* **48** 1030–69
- [54] Nethravathi C, Rajamathi J T, Ravishankar N, Shivakumara C and Rajamathi M 2008 Graphite oxide-intercalated anionic clay and its decomposition to graphene-inorganic material nanocomposites *Langmuir* **24** 8240–4
- [55] Nethravathi C, Viswanath B, Shivakumara C, Mahadevaiah N and Rajamathi M 2008 The production of smectite clay/graphene composites through delamination and co-stacking *Carbon* **46** 1773–81
- [56] Dikin D A *et al* 2007 Preparation and characterization of graphene oxide paper *Nature* **448** 457–60
- [57] Park S, Lee K, Bozkulu G, Cai W, Nguyen ST and Ruoff R S 2008 Graphene oxide papers modified by divalent ions—enhancing mechanical properties via chemical cross-linking *ACS Nano* **2** 572–8
- [58] Cote L J, Cruz-Silva R and Huang J 2009 Flash reduction and patterning of graphite oxide and its polymer composite *J. Am. Chem. Soc.* **131** 11027–32
- [59] Kim M C, Hwang G S and Ruoff R S 2009 Epoxide reduction with hydrazine on graphene: a first principles study *J. Chem. Phys.* **131** 064704
- [60] Gao X, Jang J and Nagase S 2010 Hydrazine and thermal reduction of graphene oxide: reaction mechanisms, product structures, and reaction design *J. Phys. Chem. C* **114** 832–42
- [61] Bourlino A B, Gournis D, Petridis D, Szabo T, Szeri A and Dekany I 2003 Graphite oxide: chemical reduction to graphite and surface modification with primary aliphatic amines and amino acids *Langmuir* **19** 6050–55
- [62] Gilje S, Han S, Wang M, Wang K L and Kaner R B 2007 A chemical route to graphene for device applications *Nano Lett.* **7** 3394–8
- [63] Stankovich S *et al* 2007 Synthesis of graphene-based nanosheets via chemical reduction of exfoliated graphite oxide *Carbon* **45** 1558–65
- [64] Eda G, Lin Y, Miller S, Chen C, Su W and Chhowalla M 2008 Transparent and conducting electrodes for organic electronics from reduced graphene oxide *Appl. Phys. Lett.* **92** 233305
- [65] Lomeda J R, Doyle C D, Kosynkin D V, Hwang W and Tour J M 2008 Diazonium functionalization of surfactant-wrapped chemically converted graphene sheets *J. Am. Chem. Soc.* **130** 16201–6
- [66] Zhou Y, Bao Q, Tang L A L, Zhong Y and Loh K P 2009 Hydrothermal dehydration for the ‘green’ reduction of exfoliated graphene oxide to graphene and demonstration of tunable optical limiting properties *Chem. Mater.* **21** 2950–6
- [67] Choucair M, Thordarson P and Stride J A 2009 Gram-scale production of graphene based on solvothermal synthesis and sonication *Nature Nanotechnol.* **4** 30–3
- [68] Wang G, Wang B, Park J, Yang J, Shen X and Yao J 2009 Synthesis of enhanced hydrophilic and hydrophobic graphene oxide nanosheets by a solvothermal method *Carbon* **47** 68–72
- [69] Wang H, Robinson J T, Li X and Dai H 2009 Solvothermal reduction of chemically exfoliated graphene sheets *J. Am. Chem. Soc.* **131** 9910–11
- [70] Fan X *et al* 2008 Deoxygenation of exfoliated graphite oxide under alkaline conditions: a green route to graphene preparation *Adv. Mater.* **20** 4490–3
- [71] Liu N, Luo F, Wu H, Liu Y, Zhang C and Chen J 2008 One-step ionic-liquid-assisted electrochemical synthesis of ionic-liquid-functionalized graphene sheets directly from graphite *Adv. Funct. Mater.* **18** 1518–25
- [72] Ramesha G K and Sampath S 2009 Electrochemical reduction of oriented graphene oxide films: an *in situ* Raman spectroelectrochemical study *J. Phys. Chem. C* **113** 7985–89
- [73] Zhou M *et al* 2009 Controlled synthesis of large-area and patterned electrochemically reduced graphene oxide films *Chem. Eur. J.* **15** 6116–20
- [74] Wang Z, Zhou X, Zhang J, Boey F and Zhang H 2009 Direct electrochemical reduction of single-layer graphene oxide and subsequent functionalization with glucose oxidase *J. Phys. Chem. C* **113** 14071–5
- [75] Williams G, Seger B and Kamat P V 2008 TiO₂-graphene nanocomposites. UV-assisted photocatalytic reduction of graphene oxide *ACS Nano* **2** 1487–91
- [76] Li F *et al* 2009 One-step synthesis of graphene/SnO₂ nanocomposites and its application in electrochemical supercapacitors *Nanotechnology* **20** 455602
- [77] Hernandez Y *et al* 2008 High-yield production of graphene by liquid-phase exfoliation of graphite *Nature Nanotechnol.* **3** 563–8
- [78] Geim A K and Novoselov K S 2007 The rise of graphene *Nature Mater.* **6** 183–91
- [79] Novoselov K S *et al* 2005 Two-dimensional atomic crystals *Proc. Natl Acad. Sci. USA* **102** 10451–3
- [80] Vallés C *et al* 2008 Solutions of negatively charged graphene sheets and ribbons *J. Am. Chem. Soc.* **130** 15802–4
- [81] Penicaud A, Poulin P, Derre A, Anglaret E and Petit P 2005 Spontaneous dissolution of a single-wall carbon nanotube salt *J. Am. Chem. Soc.* **127** 8–9
- [82] Bourlino A B, Georgakilas V, Zboril R, Steriotis T A and Stubos A K 2009 Liquid-phase exfoliation of graphite towards solubilized graphenes *Small* **5** 1841–5
- [83] Züchner K, Richardson T J, Glemser O and Bartlett N 1980 The pentafluoropyridine cation *Angew. Chem. Int. Edn Engl.* **19** 944–5
- [84] Langer J, Dabkowski I, Zhang Y and Illenberger E 2008 Electron capture by pentafluoronitrobenzene and pentafluorobenzonitrile *Phys. Chem. Chem. Phys.* **10** 1523–31
- [85] Lotya M *et al* 2009 Liquid phase production of graphene by exfoliation of graphite in surfactant/water solutions *J. Am. Chem. Soc.* **131** 3611–20
- [86] Moore V C *et al* 2003 Individually suspended single-walled carbon nanotubes in various surfactants *Nano Lett.* **3** 1379–82
- [87] O’Connell M J *et al* 2002 Band gap fluorescence from individual single-walled carbon nanotubes *Science* **297** 593–6
- [88] O’Connell M J *et al* 2001 Reversible water-solubilization of single-walled carbon nanotubes by polymer wrapping *Chem. Phys. Lett.* **342** 265–71
- [89] Bergin S D *et al* 2008 Large populations of individual nanotubes in surfactant-based dispersions without the

- need for ultracentrifugation *J. Phys. Chem. C* **112** 972–7
- [90] Sun Z, Nicolosi V, Rickard D, Bergin S D, Aherne D and Coleman J N 2008 quantitative evaluation of surfactant-stabilized single-walled carbon nanotubes: dispersion quality and its correlation with zeta potential *J. Phys. Chem. C* **112** 10692–9
- [91] Li D, Muller M B, Gilje S, Kaner R B and Wallace G G 2008 Processable aqueous dispersions of graphene nanosheets *Nature Nanotechnol.* **3** 101–5
- [92] Meyer J C, Geim A K, Katsnelson M I, Novoselov K S, Booth T J and Roth S 2007 The structure of suspended graphene sheets *Nature* **446** 60–3
- [93] Compton O C and Nguyen S T 2010 Graphene oxide, highly reduced graphene oxide, and graphene: versatile building blocks for carbon-based materials *Small* **6** 711–23
- [94] Allen M J, Tung V C and Kaner R B 2010 Honeycomb carbon: a review of graphene *Chem. Rev.* **110** 132–45
- [95] Loh K P, Bao Q, Ang P K and Yang J 2010 The chemistry of graphene *J. Mater. Chem.* **20** 2277